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UTILITY PATENT APPLICATION TRANSMITTAL <small>(Only for new nonprovisional applications under 37 CFR 1.53(b))</small>	Attorney Docket No. 4085-226-27 First Inventor or Application Identifier Michelle A.J. PALMER, et al Title RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME MUTANT COMPLEMENTATION
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APPLICATION ELEMENTS See MPEP chapter 600 concerning utility patent application contents		ADDRESS TO: Assistant Commissioner for Patents Box Patent Application Washington, DC 20231	
1. <input checked="" type="checkbox"/> Fee Transmittal Form (e.g. PTO/SB/17) (Submit an original and a duplicate for fee processing)		ACCOMPANYING APPLICATION PARTS <ul style="list-style-type: none"> 6. <input type="checkbox"/> Assignment Papers (cover sheet & document(s)) 7. <input type="checkbox"/> 37 C.F.R. §3.73(b) Statement <input type="checkbox"/> Power of Attorney 8. <input type="checkbox"/> English Translation Document (if applicable) 9. <input type="checkbox"/> Information Disclosure Statement (IDS)/PTO-1449 <input type="checkbox"/> Copies of IDS Citations 10. <input type="checkbox"/> Preliminary Amendment 11. <input checked="" type="checkbox"/> White Advance Serial No. Postcard 12. <input type="checkbox"/> Small Entity Statement(s) <input type="checkbox"/> Statement filed in prior application. Status still proper and desired. 13. <input type="checkbox"/> Certified Copy of Priority Document(s) (if foreign priority is claimed) 14. <input checked="" type="checkbox"/> Other: List of Inventors' Names and Addresses 	
2. <input checked="" type="checkbox"/> Specification Total Pages 25			
3. <input checked="" type="checkbox"/> Drawing(s) (35 U.S.C. 113) Total Sheets 66			
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<ul style="list-style-type: none"> a. <input type="checkbox"/> Newly executed (original or copy) b. <input type="checkbox"/> Copy from a prior application (37 C.F.R. §1.63(d)) (for continuation/divisional with box 15 completed) 			
<ul style="list-style-type: none"> i. <input type="checkbox"/> DELETION OF INVENTOR(S) Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §1.63(d)(2) and 1.33(b). 			
5. <input type="checkbox"/> Incorporation By Reference (usable if box 4B is checked) The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4B, is considered to be part of the disclosure of the accompanying application and is hereby incorporated by reference therein.			
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<input type="checkbox"/> This application is a <input type="checkbox"/> Continuation <input type="checkbox"/> Division <input type="checkbox"/> Continuation-in-part (CIP) of application Serial No. Filed on			
<input checked="" type="checkbox"/> This application claims priority of provisional application Serial No.			60/180,669
			Filed February 7, 2000
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Docket No. 4085-226-27

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

INVENTOR(S) Michelle A.J. PALMER, et al

SERIAL NO: New Application

FILING DATE: Herewith

FOR: RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS AND ORPHAN
RECEPTORS BY REPORTER ENZYME MUTANT COMPLEMENTATION

FEE TRANSMITTAL

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FOR	NUMBER FILED	NUMBER EXTRA	RATE	CALCULATIONS
TOTAL CLAIMS	30 - 20 =	10	× \$18 =	\$180.00
INDEPENDENT CLAIMS	8 - 3 =	5	× \$78 =	\$390.00
<input type="checkbox"/> MULTIPLE DEPENDENT CLAIMS (If applicable)			+ \$260 =	\$0.00
<input checked="" type="checkbox"/> LATE FILING OF DECLARATION			+ \$130 =	\$130.00
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			TOTAL OF ABOVE CALCULATIONS	\$1,390.00
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Respectfully submitted,

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TITLE OF THE INVENTION

RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME MUTANT COMPLEMENTATION

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BACKGROUND OF THE INVENTION

This application claims the benefit from Provisional Application Serial No. 60/180,669, filed February 7, 2000. The entirety of that provisional application is incorporated herein by reference.

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Field of the Invention

This invention relates to methods of detecting G-protein-coupled receptor (GPCR) activity, and provides methods of assaying GPCR activity and methods for screening for GPCR ligands, G-protein-coupled receptor kinase (GRK) activity, and compounds that interact with components of the GPCR regulatory process.

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The actions of many extracellular signals are mediated by the interaction of G-protein-coupled receptors (GPCRs) and guanine nucleotide-binding regulatory proteins (G-proteins). G-protein-mediated signaling systems have been identified in many divergent organisms, such as mammals and yeast. The GPCRs represent a large super family of proteins which have divergent amino acid sequences, but share common structural features, in particular, the presence of seven transmembrane helical domains. GPCRs respond to, among other extracellular signals, neurotransmitters, hormones, odorants and light. Individual GPCR types activate a particular signal transduction pathway; at least ten different signal transduction pathways are known to be activated via GPCRs. For example, the beta 2-adrenergic receptor (β 2AR) is a prototype mammalian GPCR. In response to agonist binding, β 2AR receptors activate a G-protein (Gs) which in turn stimulates adenylate cyclase activity and results in increased cyclic adenosine monophosphate (cAMP) production in the cell.

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The signaling pathway and final cellular response that result from GPCR stimulation depends on the specific class of G-protein with which the particular receptor is coupled (Hamm, "The many faces of G-Protein Signaling." J. Biol. Chem., 273:669-672 (1998)). For instance, coupling to the Gs class of G-proteins stimulates cAMP production and activation of Protein Kinase A and C pathways, whereas coupling to the Gi class of G-proteins down regulates cAMP. Other second messenger systems as calcium, phosphlipase C, and phosphatidylinositol 3 may also be utilized. As a consequence, GPCR signaling events have predominantly been measured via quantification of these second messenger products.

A common feature of GPCR physiology is desensitization and recycling of the receptor through the processes of receptor phosphorylation, endocytosis and dephosphorylation (Ferguson, et al., "G-protein-coupled receptor regulation: role of G-protein-coupled receptor kinases and arrestins." Can. J. Physiol. Pharmacol., 74:1095-1110 (1996)). Ligand-occupied GPCRs can be phosphorylated by two families of serine/threonine kinases, the G-protein-coupled receptor kinases (GRKs) and the second messenger-dependent protein kinases such as protein kinase A and protein kinase C. Phosphorylation by either class of kinases serves to down-regulate the receptor by uncoupling it from its corresponding G-protein. GRK-phosphorylation also serves to down-regulate the receptor by recruitment of a class of proteins known as the arrestins that bind the cytoplasmic domain of the receptor and promote clustering of the receptor into endocytic vesicles. Once the receptor is endocytosed, it will either be degraded in lysosomes or dephosphorylated and recycled back to the plasma membrane as fully-functional receptor.

Binding of an arrestin protein to an activated receptor has been documented as a common phenomenon for a variety of GPCRs ranging from rhodopsin to β2AR to the

neurotensin receptor (Barak, et al., "A β -arrestin/Green Fluorescent fusion protein biosensor for detecting G-Protein-Coupled Receptor Activation," J. Biol. Chem., 272:27497-500 (1997)). Consequently, monitoring arrestin interaction with a specific GPCR can be utilized as a generic tool for measuring GPCR activation. Similarly, a single G-protein and GRK also partner with a variety of receptors (Hamm, et al. (1998) and Pitcher et al., "G-Protein-Coupled Receptor Kinases," Annu. Rev. Biochem., 67:653-92 (1998)), such that these protein/protein interactions may also be monitored to determine receptor activity.

The present invention involves the use of a proprietary technology (ICASTTM, Intercistronic Complementation Analysis Screening Technology) for monitoring protein/protein interactions in GPCR signaling. The method involves using two inactive β -galactosidase mutants, each of which is fused with one of two interacting protein pairs, such as a GPCR and an arrestin. The formation of an active β -galactosidase complex is driven by interaction of the target proteins. In this system, β -galactosidase activity acts as a read out of GPCR activity. FIGURE 23 is a schematic depicting the method of the present invention.

FIGURE 23 shows two inactive mutants that become active when they interact. In addition, this technology could be used to monitor GPCR-mediated signaling pathways via other downstream signaling components such as G-proteins, GRKs or c-Src.

Many therapeutic drugs in use today target GPCRs, as they regulate vital physiological responses, including vasodilation, heart rate, bronchodilation, endocrine secretion and gut peristalsis. See, e.g., Lefkowitz et al., Annu. Rev. Biochem., 52:159 (1983). For instance, drugs targeting the highly studied GPCR, β 2AR are used in the treatment of anaphylaxis, shock hypertension, asthma and other conditions. Some of these drugs mimic

the ligand for this receptor. Other drugs act to antagonize the receptor in cases when disease arises from spontaneous activity of the receptor.

Efforts such as the Human Genome Project are identifying new GPCRs ("orphan" receptors) whose physiological roles and ligands are unknown. It is estimated that several thousand GPCRs exist in the human genome. Of the 250 GPCRs identified to date, only 150 have been associated with ligands.

SUMMARY OF THE INVENTION

A first aspect of the present invention is a method that monitors GPCR function proximally at the site of receptor activation, thus providing more information for drug discovery purposes due to fewer competing mechanisms. Activation of the GPCR is measured by a read-out for interaction of the receptor with a regulatory component such as arrestin, G-protein, GRK or other kinases, the binding of which to the receptor is dependent upon agonist occupation of the receptor. Protein/protein interaction is detected by complementation of reporter proteins such as utilized by the ICAST™ technology.

A further aspect of the present invention is a method of assessing G-protein-coupled receptor (GPCR) pathway activity under test conditions by providing a test cell that expresses a GPCR, e.g., muscarinic, adrenergic, dopamine, angiotensin or endothelin, as a fusion protein to a mutant reporter protein and interacting, i.e., G-proteins, arrestin or GRK, as a fusion protein with a complementing reporter protein. When test cells are exposed to a known agonist to the target GPCR under test conditions, activation of the GPCR will be monitored by complementation of the reporter enzyme. Increased reporter enzyme activity reflects interaction of the GPCR with its interacting protein partner.

A further aspect of the present invention is a method of assessing GPCR pathway activity in the presence of a test kinase.

A further aspect of the present invention is a method of assessing GPCR pathway activity in the presence of a test G-protein.

5 A further aspect of the present invention is a method of assessing GPCR pathway activity upon exposure of the test cell to a test ligand.

A further aspect of the present invention is a method of assessing GPCR pathway activity upon co-expression in the test cell of a second receptor.

10 A further aspect of the present invention is a method for screening for a ligand or agonists to an orphan GPCR. The ligand or agonist could be contained in natural or synthetic libraries or mixtures or could be a physical stimulus. A test cell is provided that expresses the orphan GPCR as a fusion protein with one β -galactosidase mutant and, for example, an arrestin or mutant form of arrestin as a fusion protein with another β -galactosidase mutant. The interaction of the arrestin with the orphan GPCR upon receptor activation is measured by 15 enzymatic activity of the complemented β -galactosidase. The test cell is exposed to a test compound, and an increase in β -galactosidase activity indicates the presence of a ligand or agonist.

20 A further aspect of the present invention is a method for screening a protein of interest, for example, an arrestin protein (or mutant form of the arrestin protein) for the ability to bind to a phosphorylated, or activated, GPCR. A cell is provided that expresses a GPCR and contains β -arrestin. The cell is exposed to a known GPCR agonist and then reporter enzyme activity is detected. Increased reporter enzyme activity indicates that the β -arrestin molecule can bind to phosphorylated, or activated, GPCR in the test cell.

10 *Am. J. Phys. Chem.* 1909, p. 101.

A further aspect of the present invention is a method to screen for an agonist to a specific GPCR. The agonist could be contained in natural or synthetic libraries or could be a physical stimulus. A test cell is provided that expresses a GPCR as a fusion protein with one β -galactosidase mutant and, for example, an arrestin as a fusion protein with another β -galactosidase mutant. The interaction of arrestin with the GPCR upon receptor activation is measured by enzymatic activity of the complemented β -galactosidase. The test cell is exposed to a test compound, and an increase in β -galactosidase activity indicates the presence of an agonist. The test cell may express a known GPCR or a variety of known GPCRs, or may express an unknown GPCR or a variety of unknown GPCRs. The GPCR may be, for example, an odorant GPCR or a β AR GPCR.

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A further aspect of the present invention is a method of screening a test compound for G-protein-coupled receptor (GPCR) antagonist activity. A test cell is provided that expresses a GPCR as a fusion protein with one β -galactosidase mutant and, for example, an arrestin as a fusion protein with another β -galactosidase mutant. The interaction of arrestin with the GPCR upon receptor activation is measured by enzymatic activity of the complemented β -galactosidase. The test cell is exposed to a test compound, and an increase in β -galactosidase activity indicates the presence of an agonist. The cell is exposed to a test compound and to a GPCR agonist, and reporter enzyme activity is detected. When exposure to the agonist occurs at the same time as or subsequent to exposure to the test compound, a decrease in β -galactosidase activity after exposure to the test compound indicates that the test compound has antagonist activity to the GPCR.

A further aspect of the present invention is a method of screening a sample solution for the presence of an agonist, antagonist or ligand to a G-protein-coupled receptor (GPCR).

A test cell is provided that expresses a GPCR fusion and contains, for example, a β-arrestin protein fusion. The test cell is exposed to a sample solution, and reporter enzyme activity is assessed. Changed reporter enzyme activity after exposure to the sample solution indicates the sample solution contains an agonist, antagonist or ligand for a GPCR expressed in the cell.

5 A further aspect of the present invention is a method of screening a cell for the presence of a G-protein-coupled receptor (GPCR).

A further aspect of the present invention is a method of screening a plurality of cells for those cells which contain a G-protein coupled receptor (GPCR).

10 A further aspect of the invention is a method for mapping GPCR-mediated signaling pathways. For instance, the system could be utilized to monitor interaction of c-src with β-arrestin-1 upon GPCR activation. Additionally, the system could be used to monitor protein/protein interactions involved in cross-talk between GPCR signaling pathways and other pathways such as that of the receptor tyrosine kinases or Ras/Raf.

15 A further aspect of the invention is a method for monitoring homo- or hetero-dimerization of GPCRs upon agonist or antagonist stimulation.

A further aspect of the invention is a method of screening a cell for the presence of a G-protein-coupled receptor (GPCR) responsive to a GPCR agonist. A cell is provided that contains protein partners that interact downstream in the GPCR's pathway. The protein partners are expressed as fusion proteins to the mutant, complementing enzyme and are used 20 to monitor activation of the GPCR. The cell is exposed to a GPCR agonist and then enzymatic activity of the reporter enzyme is detected. Increased reporter enzyme activity indicates that the cell contains a GPCR responsive to the agonist.

The invention is achieved by using ICAST™ protein/protein interaction screening to map signaling pathways. This technology is applicable to a variety of known and unknown GPCRs with diverse functions. They include, but are not limited to, the following sub-families of GPCRs:

- 5 (a) receptors that bind to amine-like ligands-Acetylcholine muscarinic receptor (M1 to M5), alpha and beta Adrenoceptors, Dopamine receptors (D1, D2, D3 and D4), Histamine receptors (H1 and H2), Octopamine receptor and Serotonin receptors (5HT1, 5HT2, 5HT4, 5HT5, 5HT6, 5HT7);
- 10 (b) receptors that bind to a peptide ligand-Angiotensin receptor, Bombesin receptor, Bradykinin receptor, C-C chemokine receptors (CCR1 to CCR8, and CCR10), C-X-C type Chemokine receptors (CXC-R5), Cholecystokinin type A receptor, CCK type receptors, Endothelin receptor, Neurotensin receptor, FMLP-related receptors, Somatostatin receptors (type 1 to type 5) and Opioid receptors (type D, K, M, X);
- 15 (c) receptors that bind to hormone proteins- Follic stimulating hormone receptor, Thyrotrophin receptor and Lutropin-choriogonadotropic hormone receptor;
- 20 (d) receptors that bind to neurotransmitters-substance P receptor, Substance K receptor and neuropeptide Y receptor;
- (e) Olfactory receptors-Olfactory type 1 to type 11, Gustatory and odorant receptors;
- (f) Prostanoid receptors-Prostaglandin E2 (EP1 to EP4 subtypes), Prostacyclin and Thromboxane;
- (g) receptors that bind to metabotropic substances-Metabotropic glutamate group I to group III receptors;

(h) receptors that respond to physical stimuli, such as light, or to chemical stimuli,

such as taste and smell; and

(i) orphan GPCRs-the natural ligand to the receptor is undefined.

ICAST™ provides many benefits to the screening process, including the ability to

5 monitor protein interactions in any sub-cellular compartment-membrane, cytosol and nucleus; the ability to achieve a more physiologically relevant model without requiring protein overexpression; and the ability to achieve a functional assay for receptor binding allowing high information content.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGURE 1. Cellular expression levels of β2 adrenergic receptor (β2AR) and β-arrestin-2 (βArr2) in C2 clones. Quantification of β-gal fusion protein was performed using antibodies against β-gal and purified β-gal protein in a titration curve by a standardized ELISA assay. Figure 1A shows expression levels of β2AR-βgalΔα clones (in expression vector pICAST ALC). Figure 1B shows expression levels of βArr2-βgalΔω in expression vector pICAST OMC4 for clones 9-3, -7, -9, -10, -19 and -24, or in expression vector pICAST OMN4 for clones 12-4, -9, -16, -18, -22 and -24.

FIGURE 2. Receptor β2AR activation was measured by agonist-stimulated cAMP production. C2 cells expressing pICAST ALC β2AR (clone 5) or parental cells were treated with increasing concentrations of (-)isoproterenol and 0.1mM IBMX. The quantification of cAMP level was expressed as pmol/well.

FIGURE 3. Interaction of activated receptor β 2AR and arrestin can be measured by

β -galactosidase complementation. Figure 3A shows a time course of β -galactosidase activity in response to agonist (-)isoproterenol stimulation in C2 expressing β 2AR- β gal $\Delta\alpha$ (β 2AR alone, in expression vector pICAST ALC), or C2 clones, and a pool of C2 co-expressing 5 β 2AR- β gal $\Delta\alpha$ and β Arr2- β gal $\Delta\omega$ (in expression vectors pICAST ALC and pICAST OMC).

Figure 3B shows a time course of β galactosidase activity in response to agonist (-)isoproterenol stimulation in C2 cells expressing β 2AR alone (in expression vector pICAST ALC) and C2 clones co-expressing β 2AR and β Arr1 (in expression vectors ICAST ALC and pICAST OMC). 10

FIGURE 4. Agonist dose response for interaction of β 2AR and arrestin can be

measured by β -galactosidase complementation. Figure 4A shows a dose response to agonists (-)isoproterenol and procaterol in C2 cells co-expressing pICAST ALC β 2AR and pICAST OMC β Arr2 fusion constructs. Figure 4B shows a dose response to agonists (-)isoproterenol and procaterol in C2 cells co-expressing pICAST ALC β 2AR and pICAST OMC β Arr1 15 fusion constructs.

FIGURE 5. Antagonist mediated inhibition of receptor activity can be measured by

β -galactosidase complementation in cells co-expressing β 2AR- β gal $\Delta\alpha$ and β Arr- β gal $\Delta\omega$. Figure 5A shows specific inhibition with adrenergic antagonists ICI-118,551 and propranolol of β -galactosidase activity in C2 clones co-expressing pICAST ALC β 2AR and pICAST OMC β Arr2 fusion constructs after incubation with agonist (-)isoproterenol. Figure 5B 20 shows specific inhibition of β -galactosidase activity with adrenergic antagonists ICI-118,551

and propranolol in C2 clones co-expressing pICAST ALC β 2AR and pICAST OMC β Arr1 fusion constructs in the presence of agonist (-)isoproterenol.

FIGURE 6. C2 cells expressing adenosine receptor A2a show cAMP induction in response to agonist (CGC-21680) treatment. C2 parental cells and C2 cells co-expressing 5 pICAST ALC A2aR and pICAST OMC β Arr1 as a pool or as selected clones were measured for agonist-induced cAMP response (pmol/well).

FIGURE 7. Agonist stimulated cAMP response in C2 cells co-expressing Dopamine receptor D1 (D1- β gal $\Delta\alpha$) and β -arrestin-2 (β Arr2- β gal $\Delta\omega$). The clone expressing β Arr2- β gal $\Delta\omega$ (Arr2 alone) was used as a negative control in the assay. Cells expressing D1- β gal $\Delta\alpha$ in addition to β Arr2- β gal $\Delta\omega$ responded agonist treatment (3-hydroxytyramine hydrochloride at 3 μ M). D1(PIC2) or D1(PIC3) designate D1 in expression vector pICAST ALC2 or pICAST ALC4, respectively.

FIGURE 8. Variety of mammalian cell lines can be used to generate stable cells for monitoring GPCR and arrestin interactions. FIGURE 8A, FIGURE 8B and FIGURE 8C show 15 the examples of HEK293, CHO and CHW cell lines co-expressing adrenergic receptor β 2AR and arrestin fusion proteins of β -galactosidase mutants. The β -galactosidase activity was used to monitor agonist-induced interaction of β 2AR and arrestin proteins.

FIGURE 9. Beta-gal complementation can be used to monitor β 2 adrenergic receptor 20 homo-dimerization. FIGURE 9A shows β -galactosidase activity in HEK293 clones co-expressing pICAST ALC β 2AR and pICAST OMC β 2AR. FIGURE 9B shows a cAMP response to agonist (-)isoproterenol in HEK 293 clones co-expressing pICAST ALC β 2AR

and pICAST OMC β 2AR. HEK293 parental cells were included in the assays as negative controls.

FIGURE 10A. pICAST ALC: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the β -gal $\Delta\alpha$; GS Linker, (GGGGS) n ; NeoR, neomycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 10B. Nucleotide sequence for pICAST ALC.

FIGURE 11A. pICAST ALN: Vector for expression of β -gal $\Delta\alpha$ as an N-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the β -gal $\Delta\alpha$; GS Linker, (GGGGS) n ; NeoR, neomycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 11B. Nucleotide sequence for pICAST ALN.

FIGURE 12A. pICAST OMC: Vector for expression of β -gal $\Delta\omega$ as a C-terminal fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the β -gal $\Delta\omega$; GS Linker, (GGGGS) n ; Hygro, hygromycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 12B. Nucleotide sequence for pICAST OMC.

FIGURE 13A. pICAST OMN: Vector for expression of β -gal $\Delta\omega$ as an N-terminal

fusion to the target protein. This construct contains the following features: MCS, multiple cloning site for cloning the target protein in frame with the β -gal $\Delta\omega$; GS Linker, (GGGGS)n;

5 Hygro, hygromycin resistance gene; IRES, internal ribosome entry site; ColE1ori, origin of replication for growth in E. coli; 5'MoMuLV LTR and 3'MoMuLV LTR, viral promotor and polyadenylation signals from the Moloney Murine leukemia virus.

FIGURE 13B. Nucleotide sequence for pICAST OMN.

FIGURE 14. pICAST ALC β Arr2: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal

10 fusion to β -arrestin-2. The coding sequence of human β -arrestin-2 (Genebank Accession Number: NM_004313) was cloned in frame to β -gal $\Delta\alpha$ in a pICAST ALC vector.

15 FIGURE 15. pICAST OMC β Arr2: Vector for expression of β -gal $\Delta\omega$ as a C-terminal fusion to β -arrestin-2. The coding sequence of human β -arrestin-2 (Genebank Accession Number: NM_004313) was cloned in frame to β -gal $\Delta\omega$ in a pICAST OMC vector.

FIGURE 16. pICAST ALC β Arr1: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal fusion to β -arrestin-1. The coding sequence of human β -arrestin-1 (Genebank Accession Number: NM_004041) was cloned in frame to β -gal $\Delta\alpha$ in a pICAST ALC vector.

20 FIGURE 17. pICAST OMC β Arr1: Vector for expression of β -gal $\Delta\omega$ as a C-terminal fusion to β -arrestin-1. The coding sequence of human β -arrestin-1 (Genebank Accession Number: NM_004041) was cloned in frame to β -gal $\Delta\omega$ in a pICAST OMC vector.

FIGURE 18. pICAST ALC β 2AR: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal fusion to β 2 Adrenergic Receptor. The coding sequence of human β 2 Adrenergic Receptor

(Genebank Accession Number: NM_000024) was cloned in frame to β -gal $\Delta\alpha$ in a pICAST ALC vector.

FIGURE 19. pICAST OMC β 2AR: Vector for expression of β -gal $\Delta\omega$ as a C-terminal fusion β 2 Adrenergic Receptor. The coding sequence of human β 2 Adrenergic Receptor (Genebank Accession Number: NM_000024) was cloned in frame to β -gal $\Delta\omega$ in a pICAST OMC vector.

FIGURE 20. pICAST ALC A2aR: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal fusion to Adenosine 2a Receptor. The coding sequence of human Adenosine 2a Receptor (Genebank Accession Number: NM_000675) was cloned in frame to β -gal $\Delta\alpha$ in a pICAST ALC vector.

FIGURE 21. pICAST OMC A2aR: Vector for expression of β -gal $\Delta\omega$ as a C-terminal fusion to Adenosine 2a Receptor. The coding sequence of human Adenosine 2a Receptor (Genebank Accession Number: NM_000675) was cloned in frame to β -gal $\Delta\omega$ in a pICAST OMC vector.

FIGURE 22. pICAST ALC D1: Vector for expression of β -gal $\Delta\alpha$ as a C-terminal fusion to Dopamine D1 Receptor. The coding sequence of human Dopamine D1 Receptor (Genebank Accession Number: X58987) was cloned in frame to β -gal $\Delta\alpha$ in a pICAST ALC vector.

FIGURE 23. A schematic depicting the method of the invention, which shows that two inactive mutants that become active when they interact.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

All literature and patents cited in this disclosure are incorporated herein by reference.

The present invention provides a method to interrogate GPCR function and pathways.

The G-protein-coupled superfamily continues to expand rapidly as new receptors are

5 discovered through automated sequencing of cDNA libraries or genomic DNA. It is

estimated that several thousand GPCRs may exist in the human genome, as many as 250

10 GPCRs have been cloned and only as few as 150 have been associated with ligands. The

means by which these, or newly discovered orphan receptors, will be associated with their

cognate ligands and physiological functions represents a major challenge to biological and

biomedical research. The identification of an orphan receptor generally requires an

individualized assay and a guess as to its function. The interrogation of a GPCR's signaling

behavior by introducing a replacement receptor eliminates these prerequisites because it can

be performed with and without prior knowledge of other signaling events. It is sensitive,

rapid and easily performed and should be applicable to nearly all GPCRs because the

15 majority of these receptors should desensitize by a common mechanism.

Various approaches have been used to monitor intracellular activity in response to a stimulant, e.g., enzyme-linked immunosorbent assay (ELISA); Fluorescence Imaging Plate

Reader assay (FLIPRTM, Molecular Devices Corp., Sunnyvale, CA); EVOscreenTM,

EVOTECTM, Evotec Biosystems GmbH, Hamburg, Germany; and techniques developed by

20 CELLOMICSTM, Cellomics, Inc., Pittsburgh, PA.

Germino, F.J., et al., "Screening for *in vivo* protein-protein interactions." Proc. Natl. Acad. Sci., 90(3): 933-7 (1993), discloses an *in vivo* approach for the isolation of proteins interacting with a protein of interest.

Phizicky, E.M., et al., "Protein-protein interactions: methods for detection and analysis." Microbiol. Rev., 59(1): 94-123 (1995), discloses a review of biochemical, molecular biological and genetic methods used to study protein-protein interactions.

Offermanns, et al., "G α ₁₅ and G α ₁₆ Couple a Wide Variety of Receptors to Phospholipase C." J. Biol. Chem., 270(25):15175-80 (1995), discloses that G α ₁₅ and G α ₁₆ can be activated by a wide variety of G-protein-coupled receptors. The selective coupling of an activated receptor to a distinct pattern of G-proteins is regarded as an important requirement to achieve accurate signal transduction. Id.

Barak et al., "A β -arrestin/Green Fluorescent Protein Biosensor for Detecting G Protein-coupled Receptor Activation." J. Biol. Chem., 272(44):27497-500 (1997) and U.S. Patent No. 5,891,646, disclose the use of a β -arrestin/green fluorescent fusion protein (GFP) to monitor protein translocation upon stimulation of GPCR.

The present invention involves a method for monitoring protein-protein interactions in GPCR pathways as a complete assay using ICAST™ (Intercistronic Complementation Analysis Screening Technology as disclosed in pending U.S. patent application serial no. 053,164, filed April 1, 1998, the entire contents of which are incorporated herein by reference). This invention enables an array of assays, including GPCR binding assays, to be achieved directly within the cellular environment in a rapid, non-radioactive assay format amenable to high-throughput screening. Using existing technology, assays of this type are currently performed in a non-cellular environment and require the use of radioisotopes.

The present invention combined with Tropix ICAST™ and Advanced Discovery Sciences™ technologies, e.g., ultra high-throughput screening, provide highly sensitive cell-based methods for interrogating GPCR pathways which are amendable to high-throughput

screening (HTS). These methods are an advancement over the invention disclosed in U.S. Patent 5,891,646, which relies on microscopic imaging of GPCR components as fusion with Green-fluorescent-protein. Imaging techniques are limited by low-throughput, lack of thorough quantification and low signal to noise ratios. Unlike yeast-based-2-hybrid assays used to monitor protein/protein interactions in high-throughput assays, the present invention is applicable to a variety of cells including mammalian cells, plant cells, protozoa cells such as E. coli and cells of invertebrate origin such as yeast, slime mold (*Dictyostelium*) and insects; detects interactions at the site of the receptor target or downstream target proteins rather than in the nucleus; and does not rely on indirect read-outs such as transcriptional activation. The present invention provides assays with greater physiological relevance and fewer false negatives.

DRAFT - INFORMATION CONTAINED HEREIN IS UNPUBLISHED AND CONFIDENTIAL

Advanced Discovery Sciences™ is in the business of offering custom-developed screening assays optimized for individual assay requirements and validated for automation. These assays are designed by HTS experts to deliver superior assay performance. Advanced Discovery Sciences'™ custom assay development service encompasses the design, development, optimization and transfer of high performance screening assays. Advanced Discovery Sciences™ works to design new assays or convert existing assays to ultra-sensitive luminescent assays ready for the rigors of HTS. Among some of the technologies developed by Advanced Discovery Sciences™ are the cAMP-Screen™ immunoassay system. This system provides ultrasensitive determination of cAMP levels in cell lysates. The cAMP-Screen™ assay utilizes the high-sensitivity chemiluminescent alkaline phosphatase (AP) substrate CSPD® with Sapphire-II™ luminescence enhancer.

EXAMPLE:

5 GPCR activation can be measured through monitoring the binding of ligand-activated
10 GPCR by an arrestin. In this assay system, a GPCR, e.g. β adrenergic receptor (β 2AR) and a
15 β arrestin are co-expressed in the same cell as fusion proteins with β gal mutants. As
20 illustrated in Figure 1, the β 2AR is expressed as a fusion protein with $\Delta\alpha$ form of β gal
mutant (β 2ADR $\Delta\alpha$) and the β arrestin as a fusion protein with the $\Delta\omega$ mutant of β gal (β -
25 Arr $\Delta\omega$). The two fusion proteins exist inside of a resting (or un-stimulated) cell in separate
30 compartments, i.e. membrane for GPCR and cytosol for arrestin, and they can not form an
35 active β galactosidase enzyme. When such a cell is treated with an agonist or a ligand, the
40 ligand-occupied and activated receptor will become a high affinity binding site for Arrestin.
45 The interaction between an activated β 2ADR $\Delta\alpha$ and β -Arr $\Delta\omega$ drives the β gal gal mutant
50 complementation. The enzyme activity can be measured by using an enzyme substrate,
55 which upon cleavage releases a product measurable by colorimetry, fluorescence,
60 chemiluminescence (e.g. Tropix product GalScreenTM).

15 **Experiment protocol-**

1. In the first step, the expression vectors for β 2ADR $\Delta\alpha$ and β Arr $\Delta\omega$ were
engineered in selectable retroviral vectors pICAST ALC, as described in Figure 18 and
pICAST OMC, as in Figure 15.
2. In the second step, the two expression constructs were transduced into either
20 C2C12 myoblast cells, or other mammalian cell lines, such as COS-7, CHO, A431, HEK 293,
and CHW. Following selection with antibiotic drugs, stable clones expressing both fusion

proteins at appropriate levels were selected.

3. In the last step, the cells expressing both β 2ADR $\Delta\alpha$ and β Arr2 $\Delta\omega$ were tested for response by agonist/ligand stimulated β galactosidase activity. Triplicate samples of cells were plated at 10,000 cells in 100 microliter volume into a well of 96-well culture plate. Cells were cultured for 24 hours before assay. For agonist assay (Figure 3 and 4), cells were treated with variable concentrations of agonist, for example, (-) isoproterenol, procaterol, dobutamine, terbutiline or L-L-phenylephrine for 60 min at 37 C. The induced β galatosidase activity was measured by addition of Tropix GalScreenTM substrate (Applied Biosystems) and luminescence measured in a Tropix TR717TM luminometer (Applied Biosystems). For antagonist assay (Figure 5), cells were pre-incubated for 10 min in fresh medium without serum in the presence of ICI-118,551 or propranolol followed by addition of 10 micro molar (-) isoproterenol.

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The assays of this invention, and their application and preparation have been described both generically, and by specific example. The examples are not intended as limiting. Other substituent identities, characteristics and assays will occur to those of ordinary skill in the art, without the exercise of inventive faculty. Such modifications remain within the scope of the invention, unless excluded by the express recitation of the claims advanced below.

WHAT IS CLAIMED IS:

1. A method of assessing the effect of a test condition on G-protein-coupled receptor (GPCR) pathway activity, comprising:
 - a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an interacting protein partner as a fusion to another mutant form of enzyme;
 - b) exposing the cell to a ligand for said GPCR under said test condition; and
 - c) monitoring activation of said GPCR by complementation of said reporter enzyme; wherein increased reporter enzyme activity in the cell compared to that which occurs in the absence of said test condition indicates increased GPCR interaction with its interacting protein partner compared to that which occurs in the absence of said test condition, and decreased reporter enzyme activity in the cell compared to that which occurs in the absence of said test condition indicates decreased GPCR interaction with its interacting protein partner compared to that which occurs in the absence of said test condition.
- 15 2. A method according to Claim 1, wherein the test condition is the presence in the cell of a kinase.
3. A method according to Claim 1, wherein the test condition is the presence in the cell of a G-protein.
4. A method according to Claim 1, wherein the test condition is the exposure of the cell to a compound selected from GPCR agonists and GPCR antagonists.
- 20 5. A method according to Claim 1, wherein the test condition is co-expression in the cell of a second receptor.
6. A method according to Claim 5, wherein the second receptor is a GPCR receptor.

7. A method according to Claim 5, wherein homo-dimerization of GPCR is determined.

8. A method according to Claim 5, wherein hetero-dimerization of GPCR is determined.

5 9. A method for screening a β -arrestin protein or an unidentified arrestin or arrestin-like protein or fragment and mutant form thereof for the ability to bind to activated GPCRs, comprising:

a) providing a cell that:

- i) expresses at least one GPCR as a fusion protein to a reporter enzyme; and
- ii) contains a conjugate comprising a test β -arrestin protein as a fusion protein

10 with another reporter enzyme;

b) exposing the cell to a ligand for said at least one GPCR; and

c) detecting enzymatic activity of the complemented reporter enzyme;

15 wherein an increase in enzymatic activity in the cell indicates β -arrestin protein

binding to the activated GPCR.

10. A method for screening a test compound for G-protein-coupled receptor (GPCR) agonist activity, comprising:

a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin protein as a fusion to another mutant form of enzyme;

20 b) exposing the cell to a test compound; and

c) detecting complementation of said reporter enzyme;

wherein increased reporter enzyme activity after exposure of the cell to the test compound indicates GPCR agonist activity of the test compound.

11. A method according to Claim 10, wherein the cell expresses a GPCR whose function is known.
12. A method according to Claim 10, wherein the cell expresses a GPCR whose function is unknown.
- 5 13. A method according to Claim 10, wherein the cell expresses an odorant or taste GPCR.
14. A method according to Claim 10, wherein the cell expresses a GPCR a β -adrenergic GPCR.
- 10 15. A method according to Claim 10, wherein the cell is selected from the group consisting of mammalian cells, cells of invertebrate origin, plant cells and protozoa cells.
16. A method according to Claim 10, wherein the cell endogenously expresses a GPCR.
17. A method according to Claim 10, wherein the cell has been transformed to express a GPCR not endogenously expressed by such a cell.

15 18. A method of screening a test compound for G-protein-coupled receptor (GPCR) antagonist activity, comprising:

- a) providing a cell that expresses a GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin protein as a fusion to another mutant form of enzyme;
- b) exposing the cell to said test compound;
- c) exposing the cell to an agonist for said GPCR; and
- d) detecting complementation of said reporter enzyme;

20 where exposure to the agonist occurs at the same time as, or subsequent to, exposure to the test compound, and wherein decreased reporter enzyme activity after exposure of the

cell to the test compound indicates that the test compound is an antagonist for said GPCR.

19. A method of screening a cell for the presence of a G-protein-coupled receptor (GPCR) responsive to a GPCR agonist, comprising:

a) providing a cell, said cell containing a conjugate comprising a β -arrestin protein as

5 a fusion protein with a reporter enzyme;

b) exposing the cell to a GPCR agonist; and

c) detecting enzymatic activity of the reporter enzyme;

wherein an increase in enzymatic activity after exposure of the cell to the GPCR
agonist indicates that the cell contains a GPCR responsive to said agonist.

10 20. A method of screening a plurality of cells for those cells which contain a G-
protein-coupled receptor (GPCR) responsive to a GPCR agonist, comprising:

a) providing a plurality of cells, said cells containing a conjugate comprising a
 β -arrestin protein as a fusion protein with a reporter enzyme;

b) exposing the cells to a GPCR agonist; and

c) detecting enzymatic activity of the reporter enzyme;

15 wherein an increase in enzymatic activity after exposure to the GPCR agonist
indicates β -arrestin protein binding to a GPCR, thereby indicating that the cell contains a
GPCR responsive to said GPCR agonist.

20 21. A method according to Claim 20, wherein the plurality of cells are contained in a
tissue.

22. A method according to Claim 20, wherein the plurality of cells are contained in
an organ.

23. A method according to Claim 20, wherein step (b) comprises exposing the cells to a plurality of GPCR agonists or ligand libraries.

24. A substrate having deposited thereon a plurality of cells, said cells expressing at least one GPCR as a fusion protein to one mutant form of reporter enzyme and an arrestin 5 protein as a fusion to another mutant form of enzyme.

25. A substrate according to Claim 24, wherein the substrate contains an enzyme-labile chemical group which, upon cleavage by the reporter enzyme, releases a product measurable by colorimetry, fluorescence or chemiluminescence.

26. A substrate according to Claim 24, wherein the substrate is made of a material selected from glass, plastic, ceramic, semiconductor, silica, fiber optic, diamond, biocompatible monomer and biocompatible polymer materials.

27. A method of detecting G-protein-coupled receptor (GPCR) pathway activity in a cell expressing at least one GPCR and containing β-arrestin protein as a fusion protein with a reporter enzyme; wherein said enzymatic activity indicates activation of the GPCR pathway.

15 28. A method according to Claim 27, where the cells are deposited on a substrate prior to detecting said enzymatic activity.

29. A method according to Claim 27, wherein said cell is contained in a tissue.

30. A method according to Claim 27, wherein said cell is contained in an organ.

ABSTRACT

Methods for detecting G-protein coupled receptor (GPCR) activity; methods of assaying GPCR activity; and methods of screening for GPCR ligands, G-protein-coupled receptor kinase (GRK) activity, and compounds that interact with components of the GPRC regulatory process are described.

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Cellular Expression of β_2 AR- β gal $\Delta\alpha$ Fusion Protein in C2 Clones
(measured by anti- β -gal ELISA)

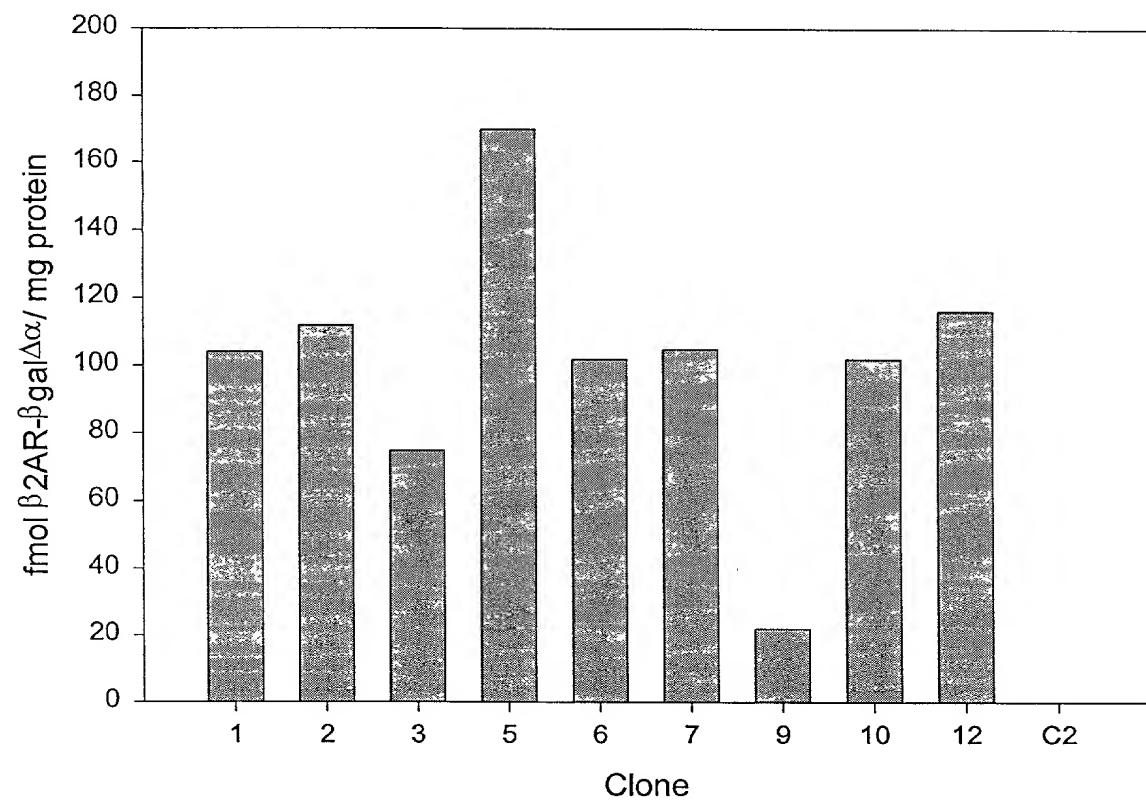


FIGURE 1A

Cellular expression of β Arr2- β gal $\Delta\omega$ fusion protein in C2 clones
(measured by anti- β gal ELISA)

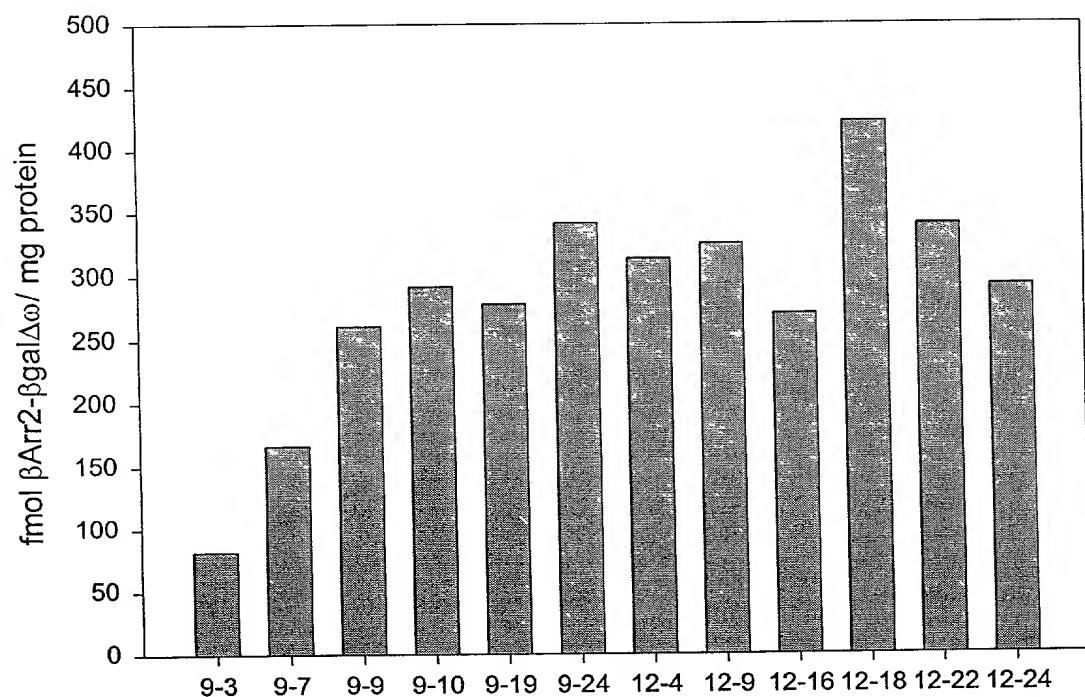


FIGURE 1B

Agonist Stimulated cAMP Response in C2 Cells Expressing β 2AR- β gal $\Delta\alpha$

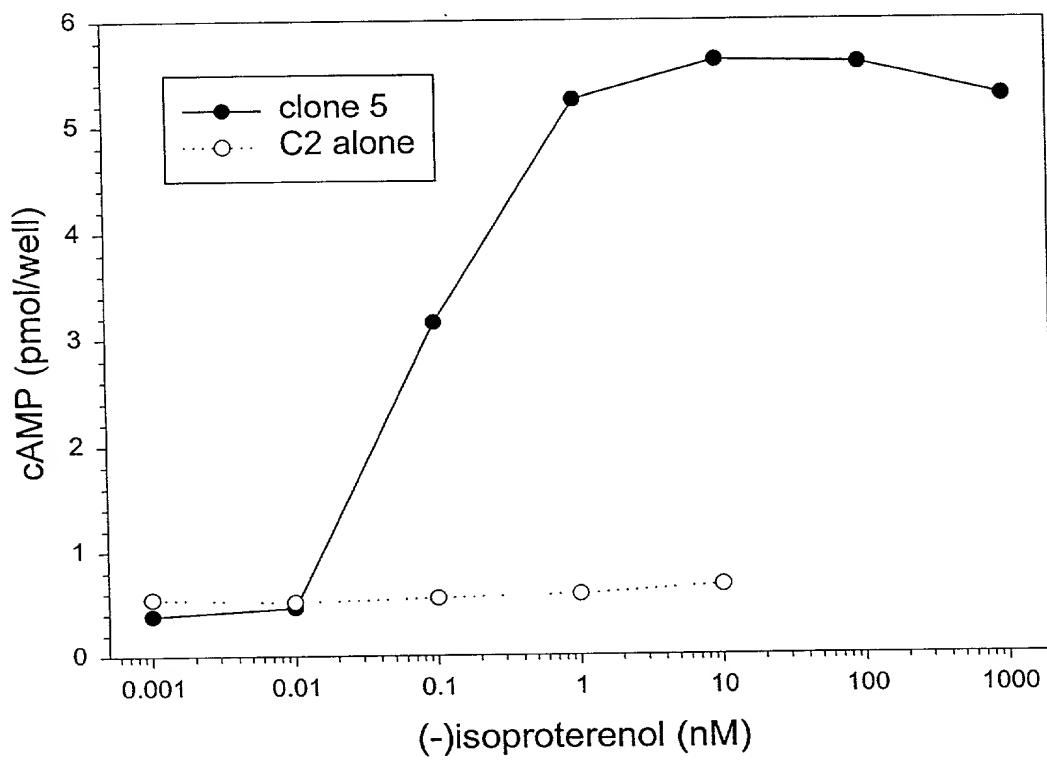


FIGURE 2

β -galactosidase Complementation as a Measurement for $\beta 2AR-\beta gal\Delta\alpha$ interacting with $\beta Arrestin2-\beta gal\Delta\omega$ upon agonist Stimulation

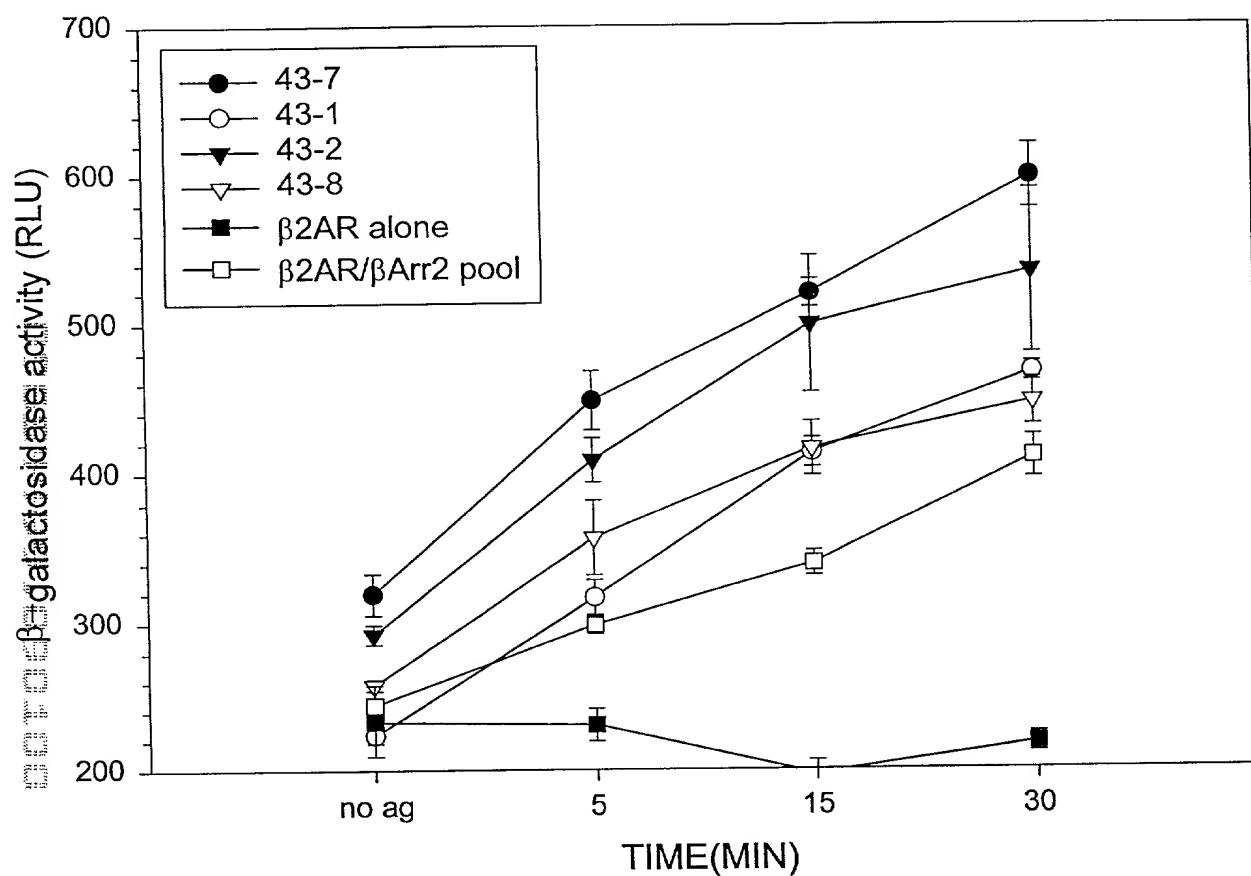


FIGURE 3A

β -galactosidase Complementation as a Measurement for $\beta 2AR-\beta gal\Delta\alpha$ Interaction with β Arrestin1- $\beta gal\Delta\omega$ upon Agonist Stimulation

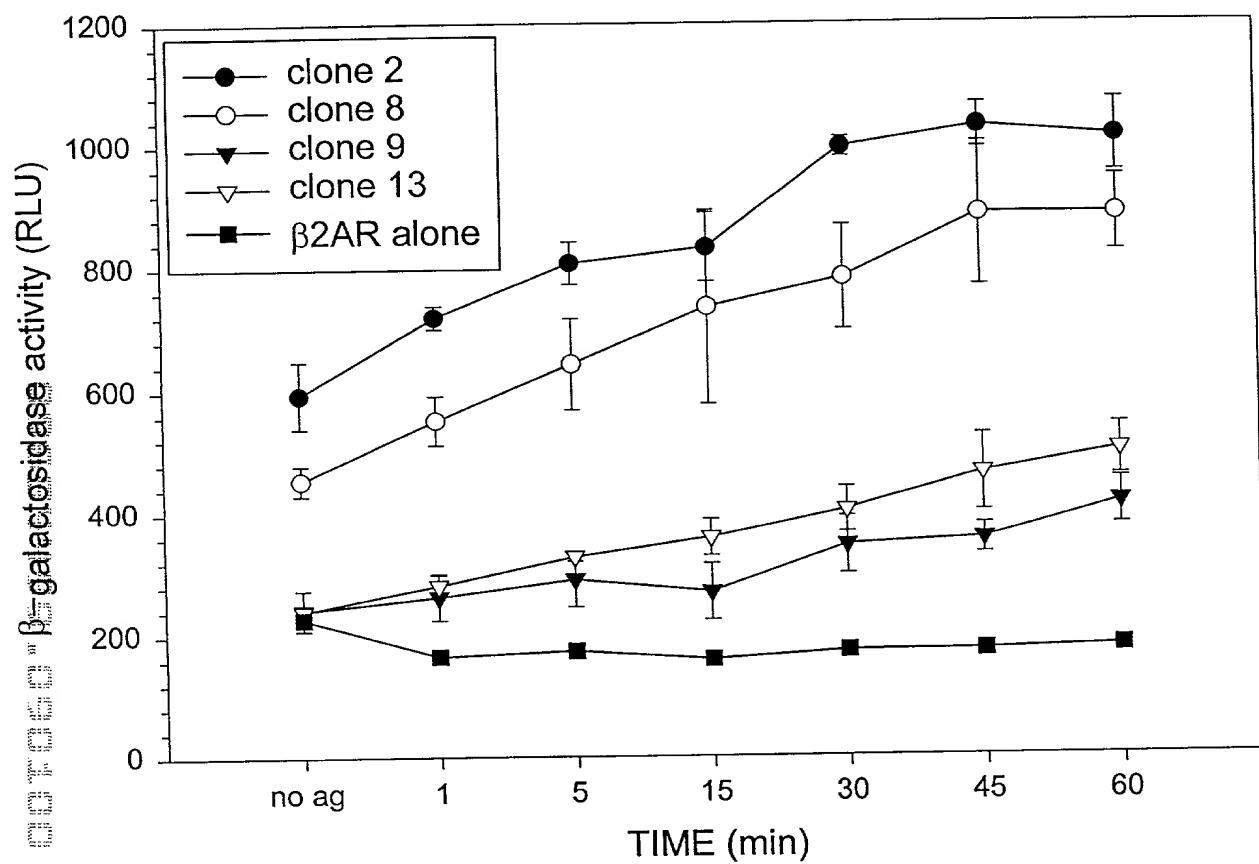


FIGURE 3B

β -galactosidase Activity in Response to Agonist in C2 Cells
Coexpressing $\beta 2\text{AR}$ - $\beta\text{gal}\Delta\alpha$ and $\beta\text{Arrestin}2$ - $\beta\text{gal}\Delta\omega$ Fusion Proteins

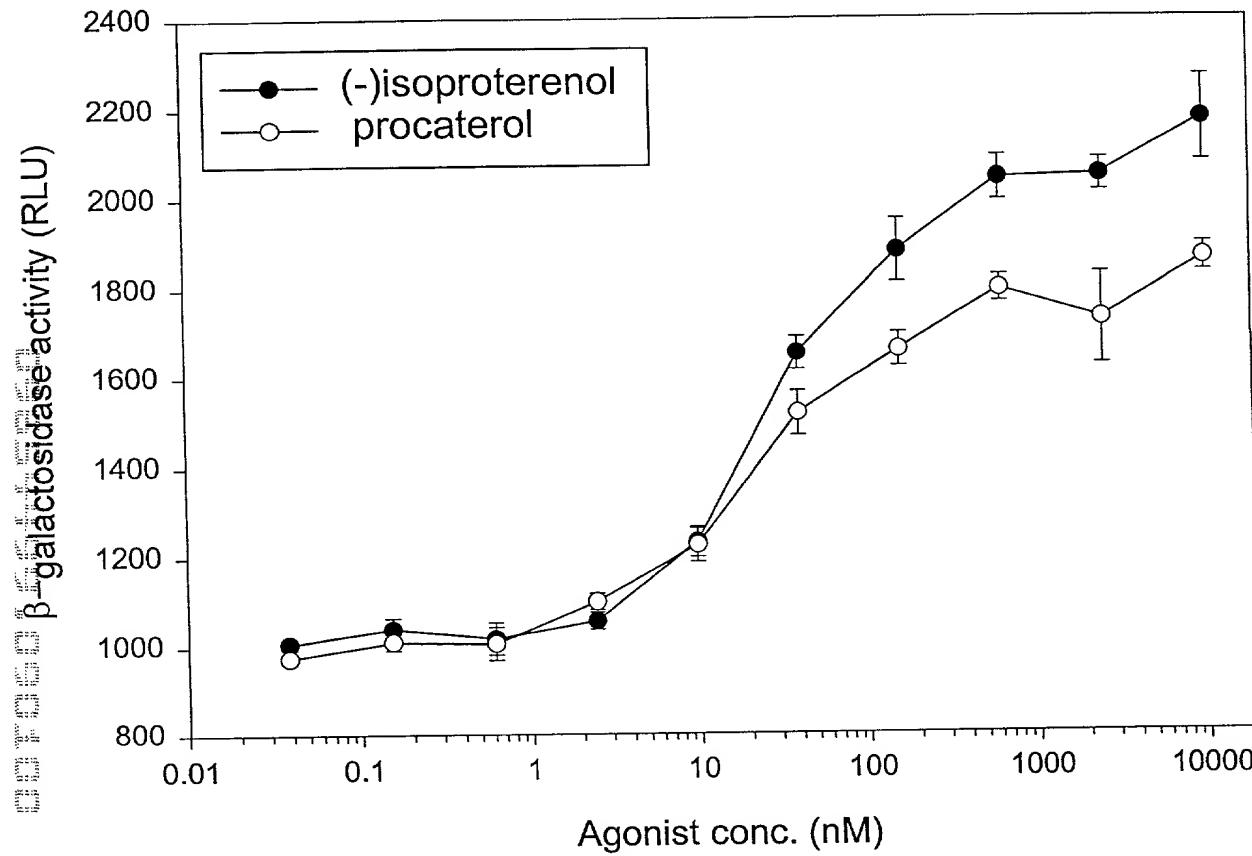


FIGURE 4A

β -galactosidase Activity in Response to Agonist in C2 Cells
Coexpressing β 2AR- β gal $\Delta\alpha$ and β Arrestin1- β gal $\Delta\omega$ Fusion Proteins

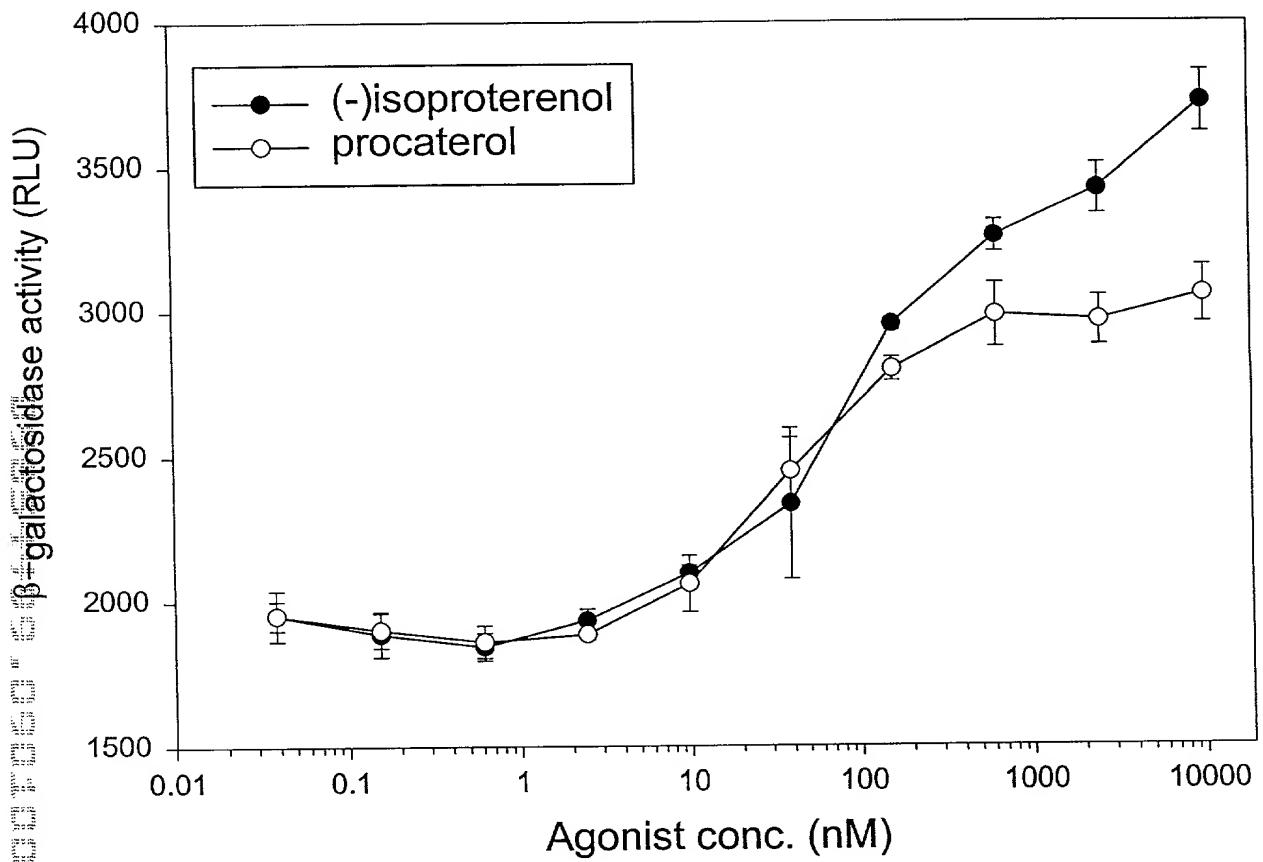


FIGURE 4B

Inhibition of β -galactosidase activity in C2 Cells Coexpressing
 $\beta 2AR-\beta gal\Delta\alpha$ and $\beta Arrestin2-\beta gal\Delta\omega$ Fusion Proteins

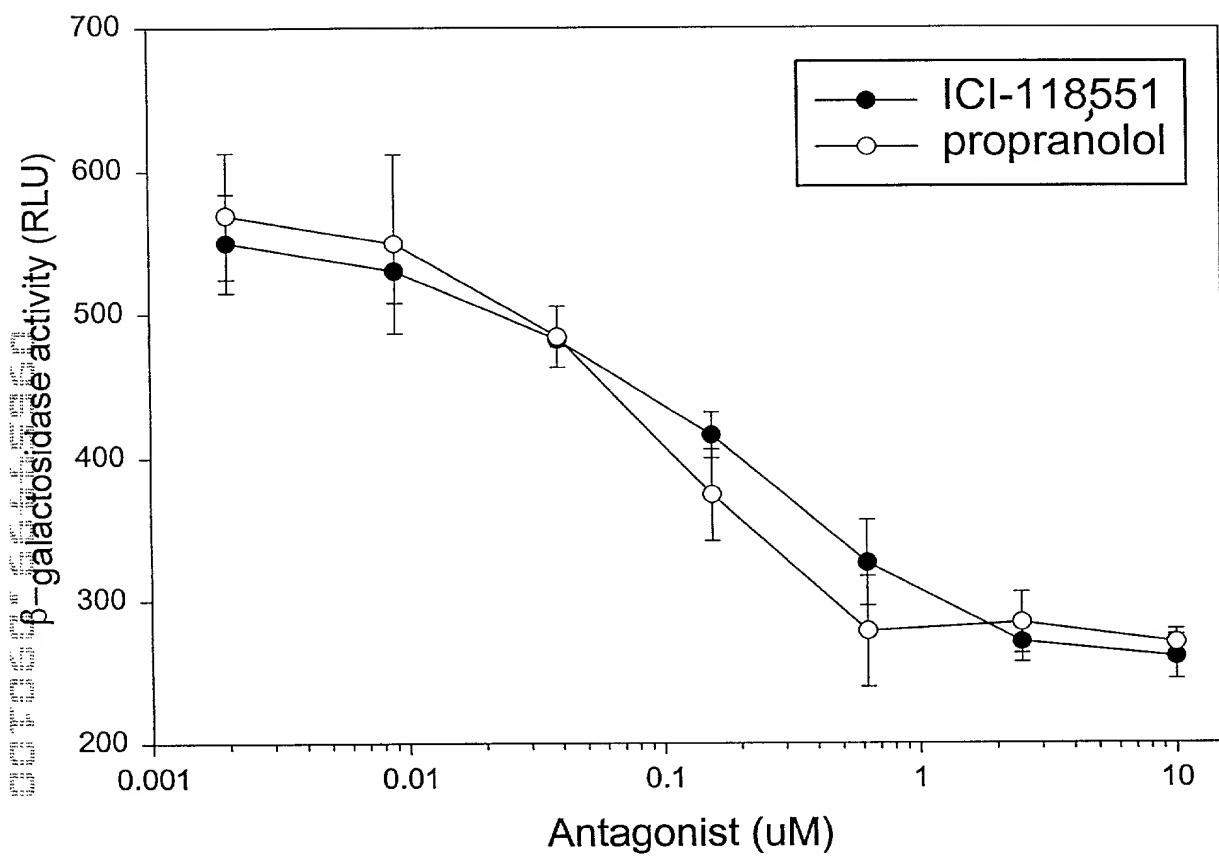


FIGURE 5A

Antagonist Inhibition of β -galactosidase Activity in C2 Cells
Coexpressing β 2AR- β gal $\Delta\alpha$ and β Arrestin1- β gal $\Delta\omega$ Fusion Proteins

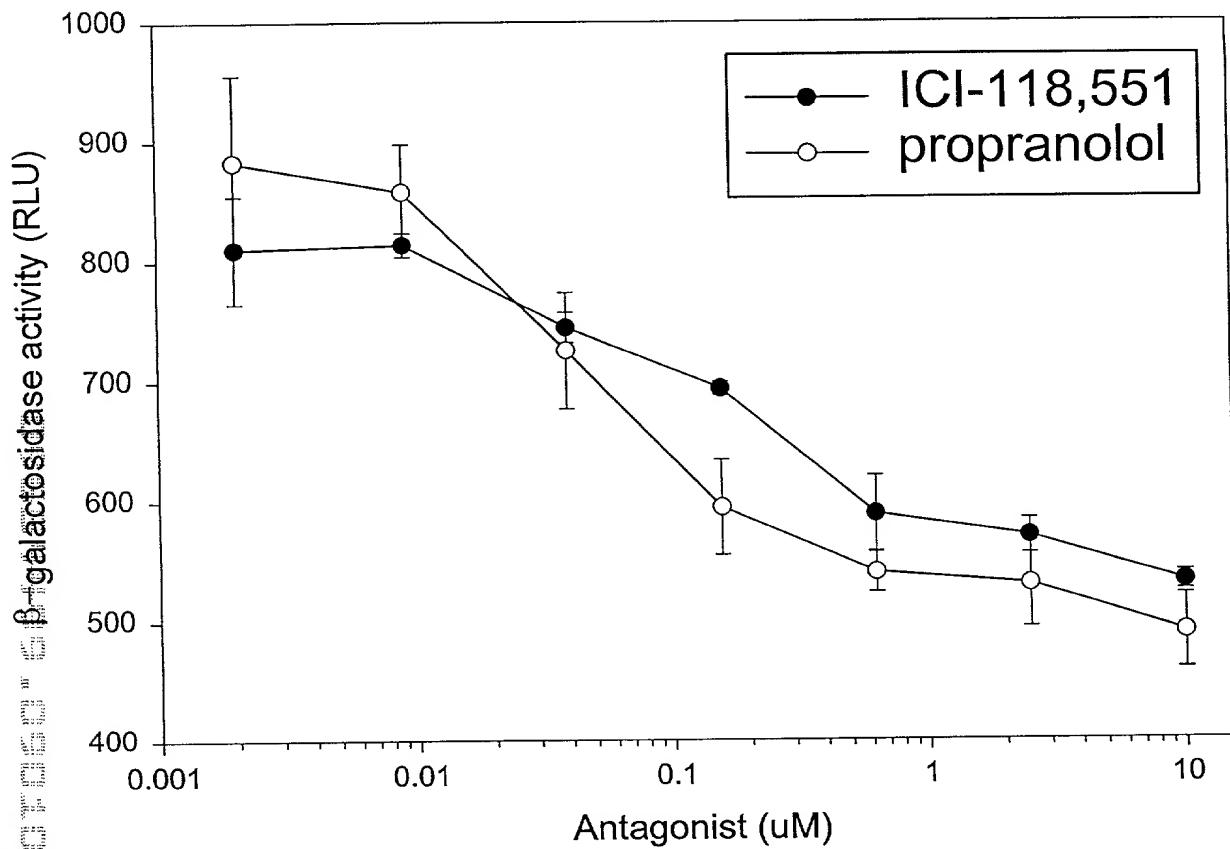


Figure 5B

Agonist Stimulated cAMP Response in Clones or Pools of C2 Cells Coexpressing A2aR- β gal $\Delta\alpha$ and β Arrestin1- β gal $\Delta\omega$ Fusion Proteins

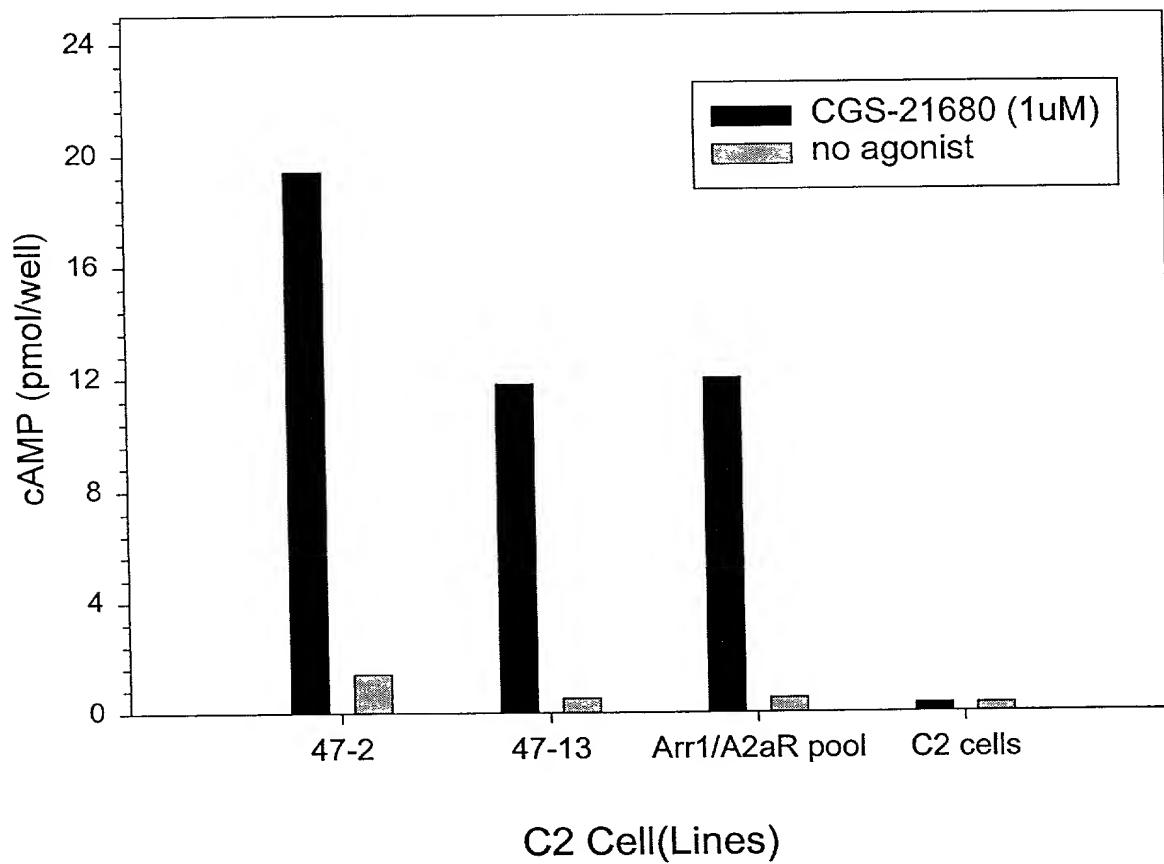


FIGURE 6

Agonist Stimulated cAMP Response in Clones or Pools of C2 Cells Expressing D1- β gal $\Delta\alpha$ and β Arrestin2- β gal $\Delta\omega$ Fusion Proteins

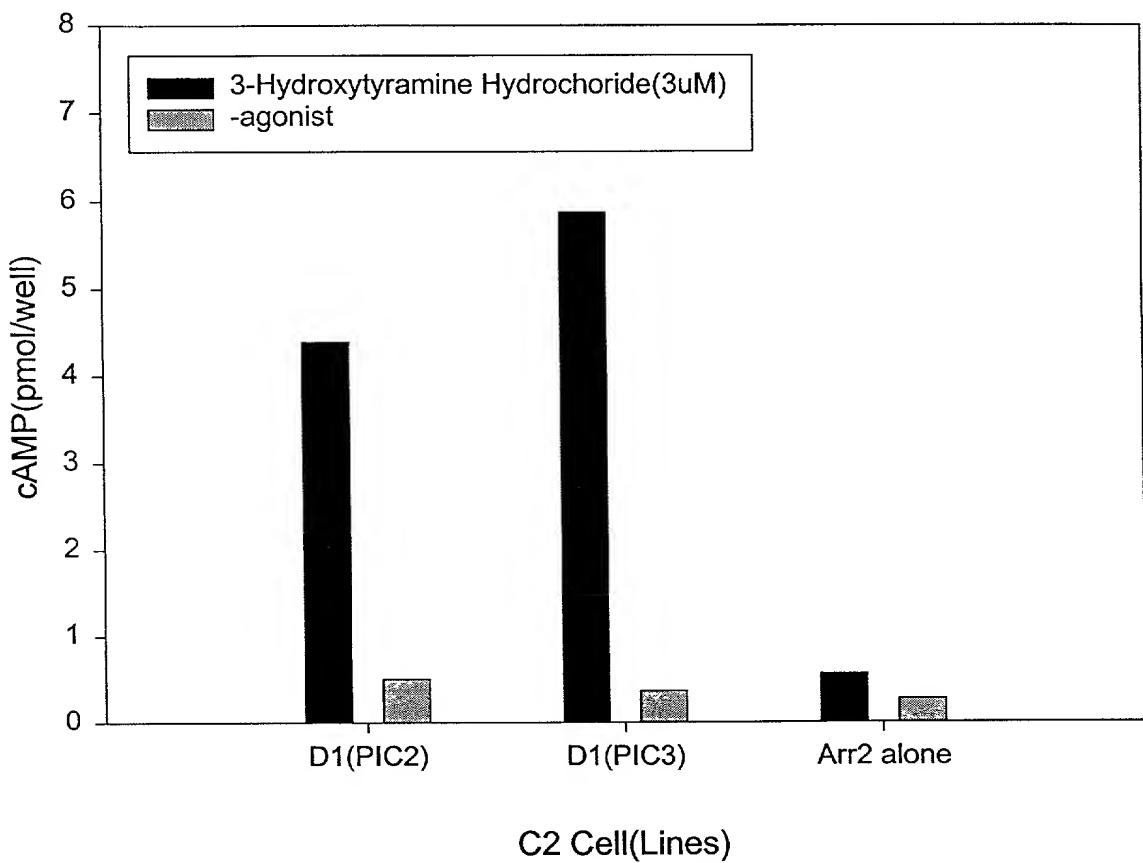


FIGURE 7

β_2 AR- β gal $\Delta\omega$ and β arr2- β gal $\Delta\alpha$ Interaction in HEK293 Clones in Response to Isoproterenol Treatment (1 μ M)

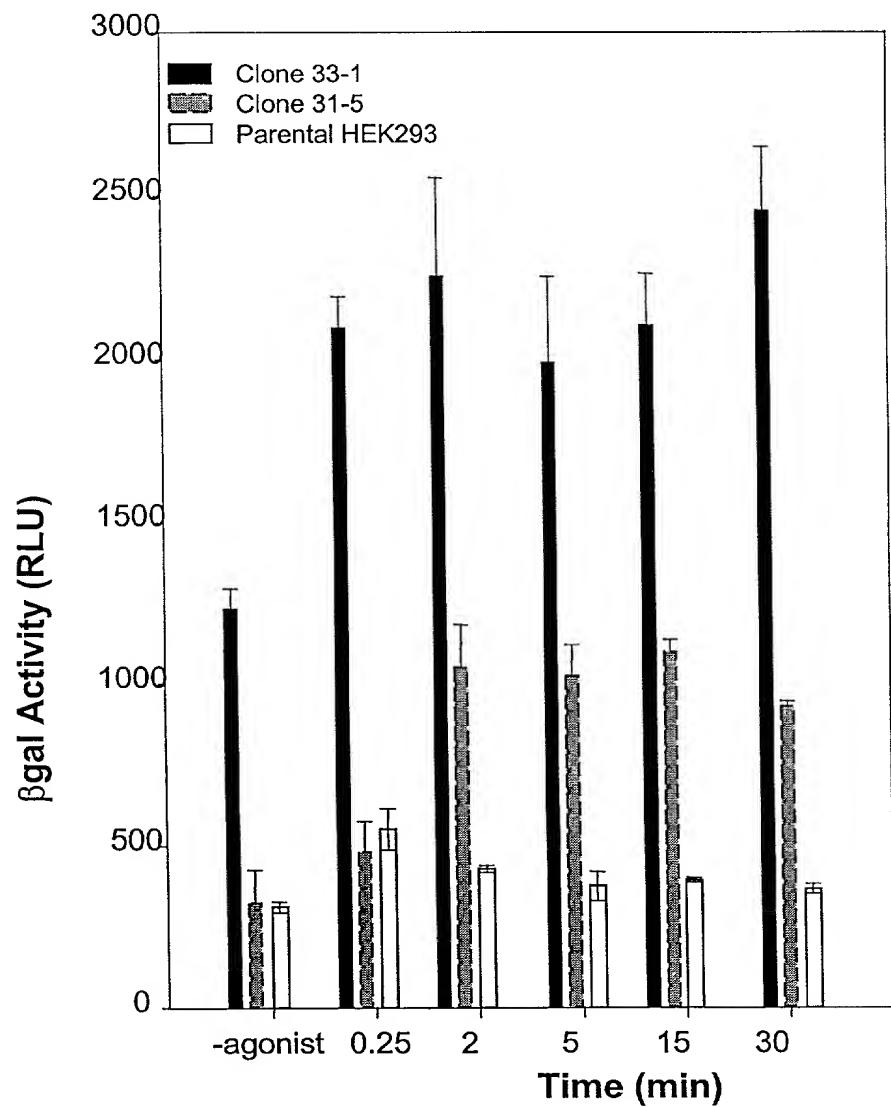


FIGURE 8A

β 2AR- β gal $\Delta\alpha$ and β Arr1- β gal Δ Interaction in a CHO Pool
in Response to Isoproterenol Treatment(10uM)

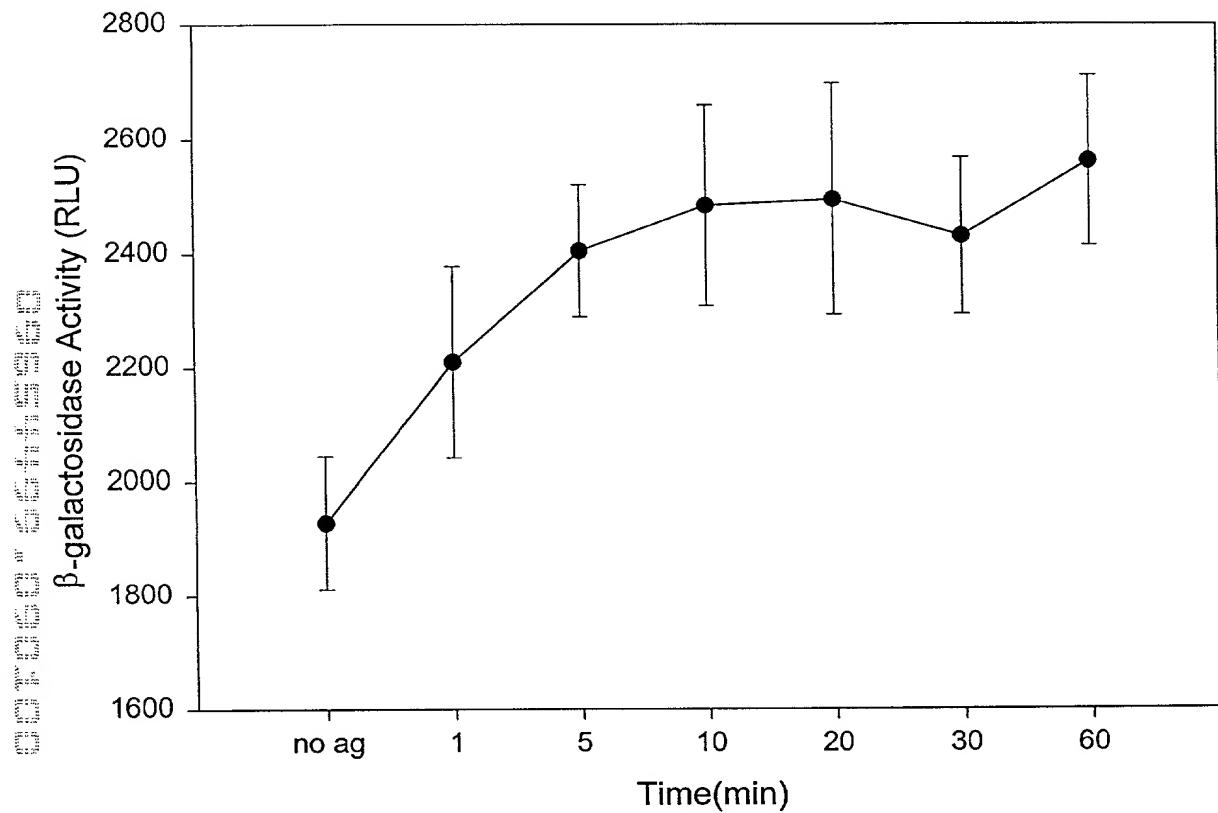


FIGURE 8B

β 2AR- β gal $\Delta\alpha$ and β Arr2- β gal $\Delta\omega$ Interaction in CHW Clone
in Response to Isoproterenol Treatment (10uM)

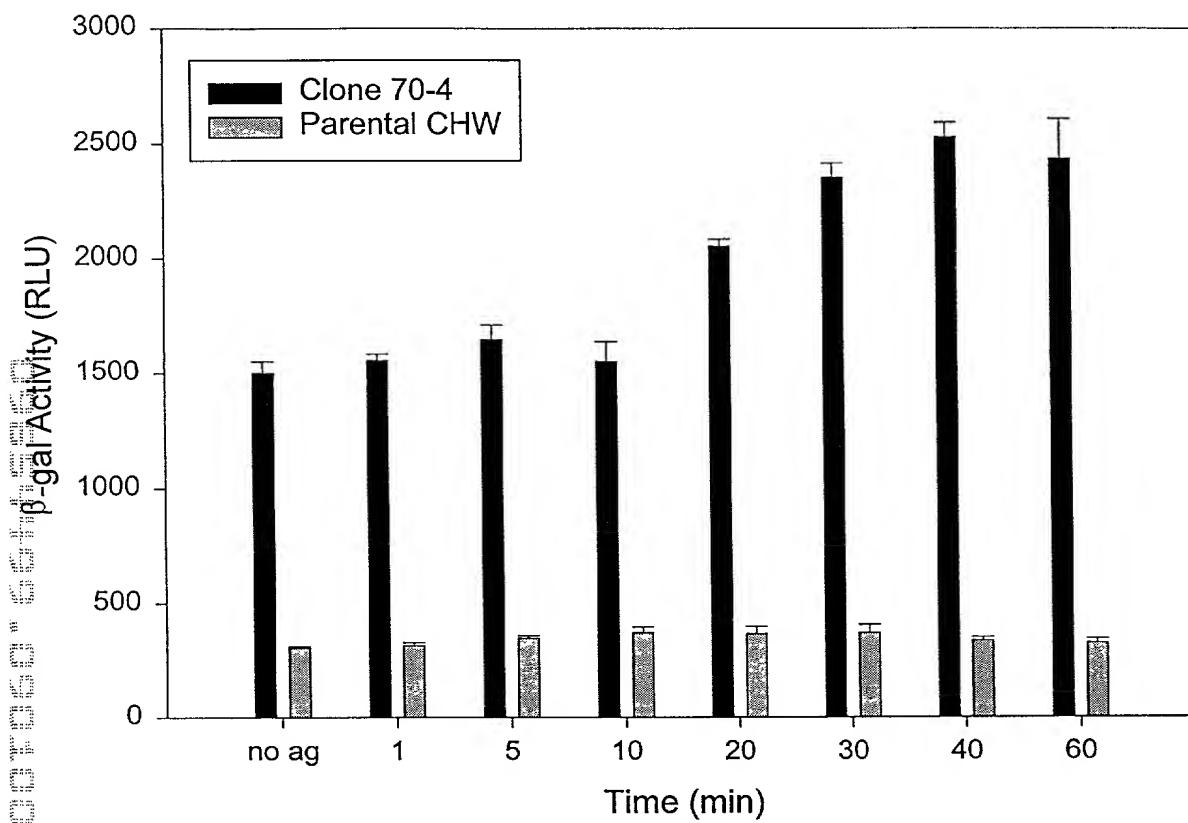


FIGURE 8C

β -galactosidase Complementation as a Measurement for Adrenergic Receptor Homodimerization in HEK 293 Cells
Coexpressing $\beta 2\text{AR}$ - $\beta\text{gal}\Delta\alpha$ and $\beta 2\text{AR}$ - $\beta\text{gal}\Delta\omega$.

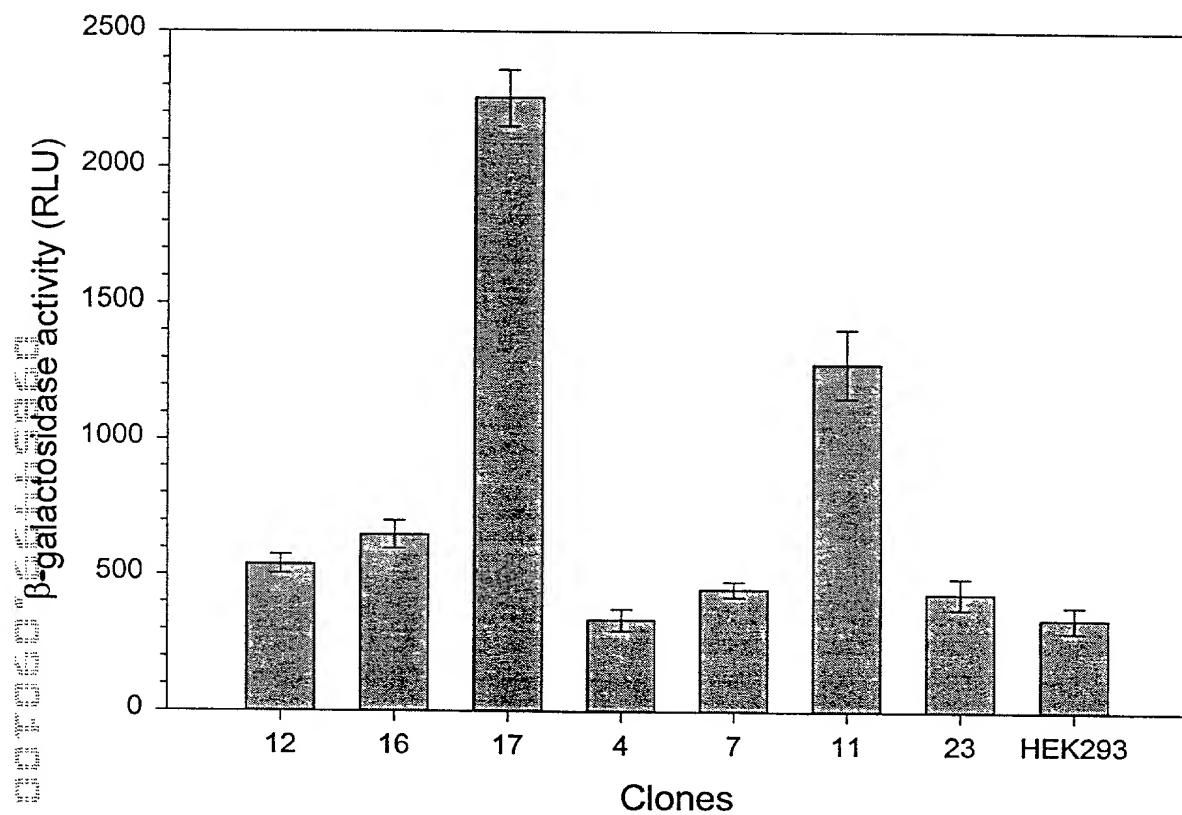


FIGURE 9A

Agonist Stimulated cAMP Response in HEK 293 Cells
Coexpressing β 2AR- β gal $\Delta\alpha$ and β 2AR- β gal $\Delta\omega$

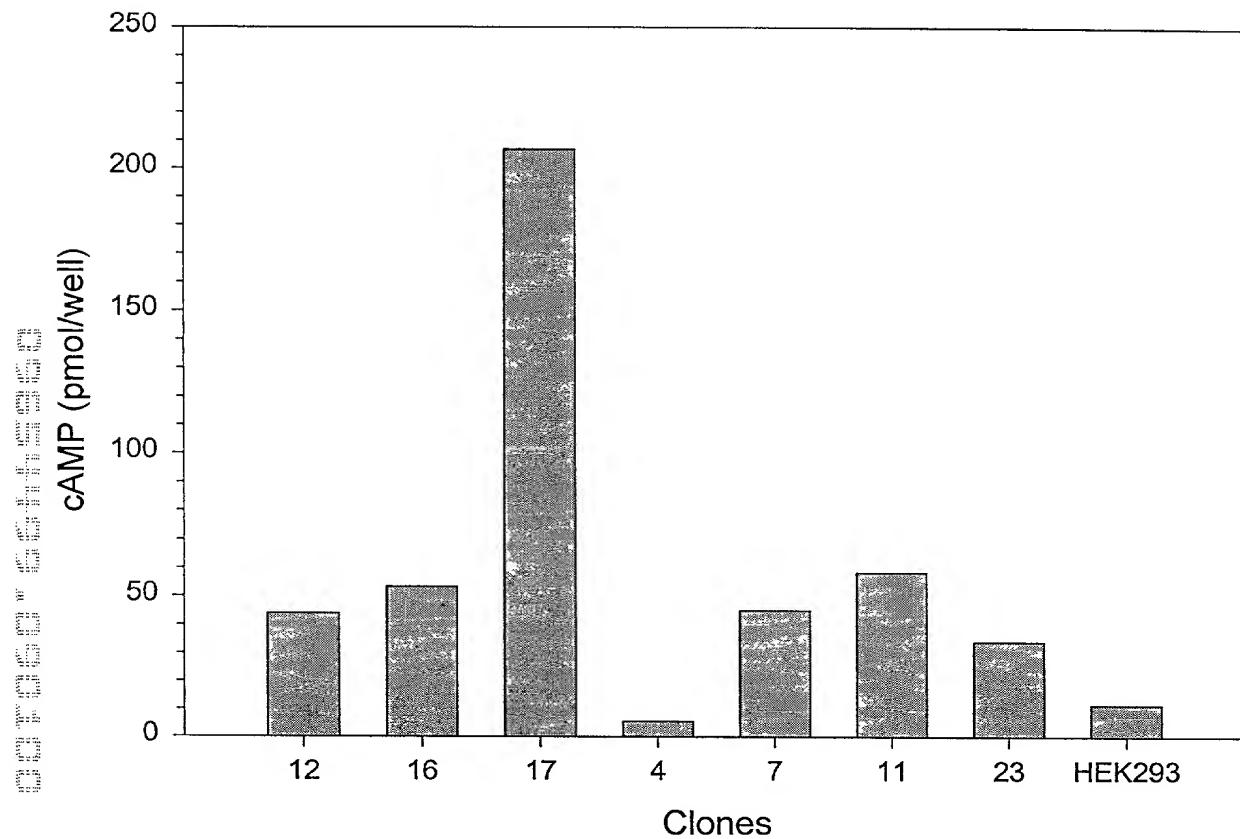


FIGURE 9B

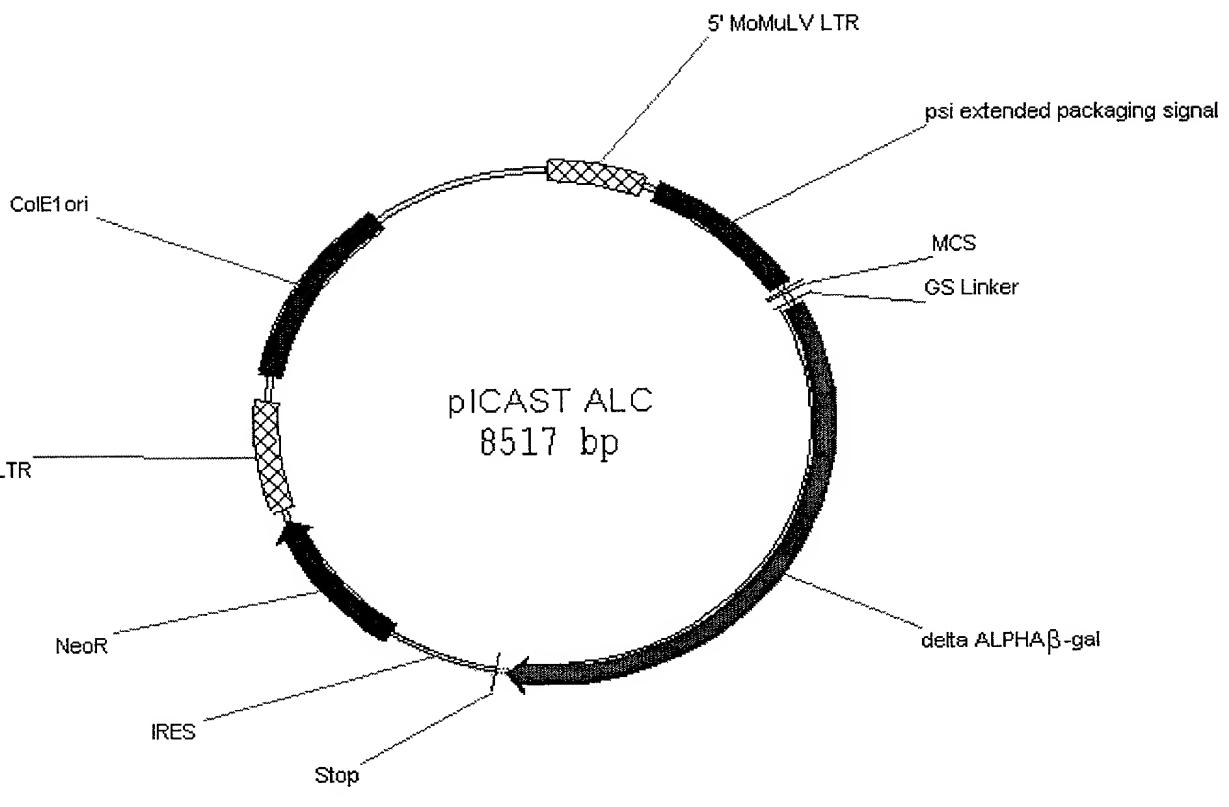


Figure 10A

1 CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
GACGTCGGAC TTATACCCGG TTTGTCCTAT AGACACCATT CGTCAAGGAC

51 CCCCCGGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTGT

101 GGATATCTGT GGTAAGCAGT TCCTGCCCCG GCTCAGGGCC AAGAACAGAT
CCTATAGACA CCATTGTCAGA AGGACGGGGC CGAGTCCCAGG TTCTTGTCTA

151 GGTCCCCAGA TCGGGTCCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA

201 GTTTCCAGGG TGCCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
CAAAGGTCCC ACGGGGTTCC TGGACTTTAC TGGGACACGG AATAAAACTTG

251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGGCT

301 GCTCAATAAA AGAGCCCACA ACCCCTCACT CGGGGGGCCA GTCCCTCCGAT
CGAGTTATT TCTCGGGTGT TGGGGAGTGA GCCCCCGGGT CAGGAGGGCTA

351 TGACTGAGTC GCCCGGGTAC CCGTGTATCC AATAAAACCTT CTTGCAGTTG
ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC

401 CATCCGACTT GTGGTCTCGC TGTTCCCTGG GAGGGTCTCC TCTGAGTGAT
GTAGGCTGAA CACCAAGAGCG ACAAGGAACC CTCCCAGAGG AGACTCACTA

451 TGACTACCCG TCAGGGGGG TCTTCATTG GGGGGCTCGT CGGGGATCGG
ACTGATGGGC AGTCGCCCCC AGAAAGTAAA CCCCCGAGCA GGCCCTAGCC

501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
CTCTGGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCAGG

551 AGCAACTTAT CTGTGTCTGT CCGATTGCTCT AGTGTCTATG ACTGATTTTA
TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAAC TGACTAAAAT

601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
ACCGGGACCG AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG

651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGCGCAACC CTGGGAGACG
GCACCACCTT GACTGCTCAA GACTTGTGGG CGGGCGTTGG GACCCTCTGC

701 TCCCAGGGAC TTTGGGGGCC GTTTTTGTGG CCCGACCTGA GGAAGGGAGT
AGGGTCCCTG AAACCCCCGG CAAAAACACC GGGCTGGACT CCTTCCCTCA

751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTCTGGT AGGAGACGAG
GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC

801 AACCTAAAAC AGTTCCCGCC TCCGTCTGAA TTTTGCTTT CGGTTGGAA
TTGGATTGG TCAAGGGCGG AGGCAGACTT AAAAACGAAA GCCAAACCTT

851 CCGAAGCCGC GCGTCTTGTG TGCTGCAGCA TCGTTCTGTG TTGTCTCTGT
GGCTTCGGCG CGCAGAACAG ACAGACGTCGT AGCAAGACAC AACAGAGACA

901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAAC
GACTGACACA AAGACATAAA CAGACTTTA ATCCCGGTCT GACAATGGTG

FIGURE 10B

951 TCCCTTAAGT TTGACCTTAG GTAACTGGAA AGATGTCGAG CGGCTCGCTC
AGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG

1001 ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
TGTTGGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA

1051 GCAGAACATGGC CAACCTTTAA CGTCGGATGG CCGCGAGACG GCACCTTTAA
CGTCTTACCG GTTGGAAATT GCAGCCTACC GGCGCTCTGC CGTGGAAATT

1101 CCGAGACCTC ATCACCCAGG TTAAGATCAA GGTCTTTCA CCTGGCCCCGC
GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAAGT GGACCGGGCG

1151 ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCCT TCGGAACCGA

1201 TTTGACCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCCTA AGCCTCCGCC
AAACTGGGGG GAGGGACCCA GTTCGGGAAA CATGTGGGAT TCGGAGGCAG

1251 TCCTCTTCTT CCATCCGCC CGTCTCTCCC CCTTGAACCT CCTCGTTCGA
AGGAGAAGGA GGTAGGGCGGG GCAGAGAGGG GGAACCTGGGA GGAGCAAGCT

1301 CCCC CGCCTCG ATCCTCCCTT TATCCAGCCC TCACCTCTTC TCTAGGCGCC
GGGGCGGAGC TAGGAGGGAA ATAGGTCGGG AGTGAGGAAG AGATCCGCAG

1351 GGCCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAATCAGG
CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCGCTA AGCTTAGTCC

1401 CCTTGGCGCG CGGGATCCTT AATTAAGCGC AATTGGGAGG TGGCGGTAGC
GGAACCGCGC GGCCTAGGAA TTAATTGCGC TTAACCCCTCC ACCGCCATCG

+2 M G V I T D S L A V V A R T D

1451 CTCGAGATGG GCGTGATTAC GGATTCACTG GCCGTCGTGG CCCGCACCGA
GAGCTCTACC CGCACTAATG CCTAAGTGAC CGGCAGCACC GGGCGTGGCT

+2 R P S Q Q L R S L N G E W R F A

1501 TCGCCCTTCC CAACAGTTAC GCAGCCTGAA TGGCGAATGG CGCTTTGCCT
AGCGGGAAAGG GTTGTCAATG CGTCGGACTT ACCGCTTACC GCGAAACGGA

+2 W F P A P E A V P E S W L E C D L

1551 GGTTTCCGGC ACCAGAAGCG GTGCCGGAAA GCTGGCTGGA GTGCGATCTT
CCAAAGGCCG TGGTCTTCGC CACGGCCTTT CGACCGACCT CACGCTAGAA

+2 P E A D T V V V P S N W Q M H G Y

1601 CCTGAGGCCG ATACTGTCGT CGTCCCCCTCA AACTGGCAGA TGACCGGTTA
GGACTCCGGC TATGACAGCA GCAGGGGAGT TTGACCGTCT ACGTGCCAAT

+2 D A P I Y T N V T Y P I T V N P

1651 CGATGCGCCC ATCTACACCA ACGTGACCTA TCCCATTACG GTCAATCCGC
GCTACGCCGG TAGATGTGGT TGCACGGAT AGGGTAATGC CAGTTAGGCG

+2 P F V P T E N P T G C Y S L T F N

1701 CGTTTGTTC CACGGAGAAT CCGACGGGTT GTTACTCGCT CACATTAAAT
GCAAACAAGG GTGCCTCTTA GGCTGCCAA CAATGAGCGA GTGTAAATTA

+2 V D E S W L Q E G Q T R I I F D G

1751 GTTGATGAAA GCTGGCTACA GGAAGGCCAG ACGCGATTAA TTTTGATGG
CAACTACTTT CGACCGATGT CCTTCCGGTC TGCGCTTAAT AAAAACTACC

+2 V N S A F H L W C N G R W V G Y

1801 CGTTAACTCG GCGTTTCATC TGTGGTGCAA CGGGCGCTGG GTCGGTTACCG
GCAATTGAGC CGCAAAGTAG ACACCACGTT GCCCGCGACC CAGCCAATGC

+2 G Q D S R L P S E F D L S A F L R

1851 GCCAGGACAG TCGTTGCGC TCTGAATTG ACCTGAGCGC ATTTTACGC
CGGTCCCTGTC AGCAAACGGC AGACTTAAAC TGGACTCGCG TAAAAATGCG

+2 A G E N R L A V M V L R W S D G S

1901 GCCGGAGAAA ACCGCCTCGC GGTGATGGTG CTGCGCTGGA GTGACGGCAG
CGGCCTCTT TGGCGGAGCG CCACTACCAC GACGCGACCT CACTGCCGTC

+2 Y L E D Q D M W R M S G I F R D

1951 TTATCTGGAA GATCAGGATA TGTGGCGGAT GAGCGGCATT TTCCGTGACG
AATAGACCTT CTAGTCCTAT ACACCGCCTA CTCGCCGTAA AAGGCACTGC

+2 V S L L H K P T T Q I S D F H V A

2001 TCTCGTTGCT GCATAAACCG ACTACACAAA TCAGCGATTT CCATGTTGCC
AGAGCAACGA CGTATTTGGC TGATGTGTTT AGTCGCTAAA GGTACAACGG

+2 T R F N D D F S R A V L E A E V Q

2051 ACTCGCTTTA ATGATGATTT CAGCCGCGCT GTACTGGAGG CTGAAGTTCA
TGAGCGAAAT TACTACTAAA GTCGGCGCGA CATGACCTCC GACTTCAAGT

+2 M C G E L R D Y L R V T V S L W

2101 GATGTGCGGC GAGTTGCGTG ACTACCTACG GGTAAACAGTT TCTTTATGGC
CTACACGCCG CTCAACGCAC TGATGGATGC CCATTGTCAA AGAAATACCG

+2 Q G E T Q V A S G T A P F G G E I

2151 AGGGTAAAC GCAGGTGCGC AGCGGCACCG CGCCCTTCGG CGGTGAAATT
TCCCACTTTG CGTCCAGCGG TCGCCGTGGC GCGGAAAGCC GCCACTTTAA

+2 I D E R G G Y A D R V T L R L N V

2201 ATCGATGAGC GTGGTGGTTA TGCCGATCGC GTCACACTAC GTCTGAACGT
TAGCTACTCG CACCACCAAT ACGGCTAGCG CAGTGTGATG CAGACTTGCA

+2 E N P K L W S A E I P N L Y R A

2251 CGAAAACCCG AACTGTGGA GCGCCGAAAT CCCGAATCTC TATCGTGCGG
GCTTTGGGC TTGACACCT CGCGCTTTA GGGCTTAGAG ATAGCACGCC

+2 V V E L H T A D G T L I E A E A C

2301 TGGTTGAACT GCACACCGCC GACGGCACGC TGATTGAAGC AGAAGCCTGC
ACCAACTTGA CGTGTGGCGG CTGCCGTGCG ACTAACTTCG TCTTCGGACG

+2 D V G F R E V R I E N G L L L L N

2351 GATGTCGGTT TCCCGGAGGT GCGGATTGAA AATGGTCTGC TGCTGCTGAA
CTACAGCCAA AGGCCTCCA CGCCTAACCTT TTACCAAGACG ACGACGACTT

+2 G K P L L I R G V N R H E H H P

2401 CGGCAAGCCG TTGCTGATTC GAGGCGTTAA CCGTCACGAG CATCATCCTC
GCCGTTCCGC AACGACTAAG CTCCGCAATT GGCAGTGCTC GTAGTAGGAG

+2 L H G Q V M D E Q T M V Q D I L L

2451 TGCATGGTCA GGTCACTGGAT GAGCAGACGA TGGTGCAGGA TATCCTGCTG
ACGTACCACT CCAGTACCTA CTCGTCTGCT ACCACGTCC ATAGGACGAC

+2 M K Q N N F N A V R C S H Y P N H

2501 ATGAAGCAGA ACAACTTAA CGCCGTGCGC TGTTCGCATT ATCCGAACCA
TACTTCGTCT TGTGAAATT GCGGCACGCG ACAAGCGTAA TAGGCTTGGT

+2 P L W Y T L C D R Y G L Y V V D

2551 TCCGCTGTGG TACACGCTGT GCGACCGCTA CGGCTGTAT GTGGTGGATG
AGGCGACACC ATGTGCGACA CGCTGGCGAT GCCGGACATA CACCACCTAC

+2 E A N I E T H G M V P M N R L T D

2601 AAGCCAATAT TGAAACCCAC GGCATGGTGC CAATGAATCG TCTGACCGAT
TCGGTTATA ACTTGGGTG CCGTACCAAG GTTACTTAGC AGACTGGCTA

+2 D P R W L P A M S E R V T R M V Q

2651 GATCCGGCGCT GGCTACCGGC GATGAGCGAA CGCGTAACGC GAATGGTGCA
CTAGGCGCGA CCGATGGCCG CTACTCGCTT GCGCATTGCG CTTACCACGT

+2 R D R N H P S V I I W S L G N E

2701 GCGCGATCGT AATCACCCGA GTGTGATCAT CTGGTCGCTG GGGAAATGAAT
CGCGCTAGCA TTAGTGGCT CACACTAGTA GACCAGCGAC CCCTTACTTA

+2 S G H G A N H D A L Y R W I K S V

2751 CAGGCCACGG CGCTAACAC GACCGCTGT ATCGCTGGAT CAAATCTGTC
GTCCGGTGCC GCGATTAGTG CTGCGCACA TAGGGACCTA GTTTAGACAG

+2 D P S R P V Q Y E G G G A D T T A

2801 GATCCTTCCC GCCCCGTGCA GTATGAAGGC GGCAGGAGCCG ACACCACGGC
CTAGGAAGGG CGGGCACGT CATACTTCCG CCGCCTCGGC TGTGGTGCCG

+2 T D I I C P M Y A R V D E D Q P

2851 CACCGATATT ATTGCCCCGA TGTACGCGCG CGTGGATGAA GACCAGCCCT
GTGGCTATAA TAAACGGGCT ACATGCGCGC GCACCTACTT CTGGTCGGGA

+2 F P A V P K W S I K K W L S L P G

2901 TCCC GGCTGT GCCGAAATGG TCCATCAAAA AATGGCTTTC GCTACCTGGA
AGGGCCGACA CGGCTTACC AGGTAGTTT TTACCGAAAG CGATGGACCT

+2 E T R P L I L C E Y A H A M G N S

2951 GAGACGCGCC CGCTGATCCT TTGCGAATAC GCCCCACGCGA TGGGTAACAG
CTCTGCGCGG GCGACTAGGA AACGCTTATG CGGGTGCCT ACCCATTGTC

+2 L G G F A K Y W Q A F R Q Y P R

3001 TCTTGGCGGT TTCGCTAAAT ACTGGCAGGC GTTCGTCAG TATCCCCGT
AGAACCGCCA AAGCGATTTA TGACCGTCCG CAAAGCAGTC ATAGGGGCAA

+2 L Q G G F V W D W V D Q S L I K Y

3051 TACAGGGCGG CTTCGTCTGG GACTGGGTGG ATCAGTCGCT GATTAAATAT
ATGTCCCGCC GAAGCAGACC CTGACCCACC TAGTCAGCGA CTAATTATA

+2 D E N G N P W S A Y G G D F G D T

3101 GATGAAAACG GCAACCCGTG GTCGGCTTAC GGCGGTGATT TTGGCGATAC
CTACTTTGC CGTTGGCAC CAGCCGAATG CCGCCACTAA AACCGCTATG

+2 P N D R Q F C M N G L V F A D R

3151 GCGGAACGAT CGCCAGTTCT GTATGAACGG TCTGGTCTTT GCCGACCGCA
CGGCTTGCTA CGGGCTAACAGA CATACTTGCC AGACCAGAAA CGGCTGGCGT

+2 T P H P A L T E A K H Q Q Q F F Q

3201 CGCCGCATCC AGCGCTGACG GAAGCAAAAC ACCAGCAGCA GTTTTTCCAG
GCGGCGTAGG TCGCGACTGC CTTCGTTTG TGGTCGTCGT CAAAAAGGTC

+2 F R L S G Q T I E V T S E Y L F R

3251 TTCCGTTTAT CCGGGCAAC CATCGAAGTG ACCAGCGAAT ACCTGTTCCG
AAGGCAAATA GGCCCGTTG GTAGCTTCAC TGGTCGCTTA TGGACAAGGC

+2 H S D N E L L H W M V A L D G K

3301 TCATAGCGAT AACGAGCTCC TGCACGGAT GGTGGCGCTG GATGGTAAGC
AGTATCGCTA TTGCTCGAGG ACGTGACCTA CCACCGCGAC CTACCATTCG

+2 P L A S G E V P L D V A P Q G K Q

3351 CGCTGGCAAG CGGTGAAGTG CCTCTGGAT GCGCTCCACA AGGTAAACAG
GCGACCGTTC GCCACTTCAC GGAGACCTAC AGCGAGGTGT TCCATTTGTC

+2 L I E L P E L P Q P E S A G Q L W

3401 TTGATTGAAC TGCTGAACT ACCGCAGCCG GAGAGCGCCG GGCAACTCTG
AACTAACTTG ACGGACTTGA TGGCGTCGGC CTCTCGCGC CCGTTGAGAC

+2 L T V R V V Q P N A T A W S E A

3451 GCTCACAGTA CGCGTAGTGC AACCGAACGC GACCGCATGG TCAGAACGCC
CGAGTGTCACT GCGCATCACG TTGGCTTGCG CTGGCGTACC AGTCTTCGGC

+2 G H I S A W Q Q W R L A E N L S V

3501 GGCACATCATCG CGCCTGGCAG CAGTGGCGTC TGGCGAAAA CCTCAGTGTG
CCGTGTAGTC CGGGACCGTC GTCACCGCAG ACCGCCTTT GGACTCACAC

+2 T L P A A S H A I P H L T T S E M

3551 ACGCTCCCCG CCGCGTCCC CGCCCATCCCG CATCTGACCA CCAGCGAAAT
TGCAGGGGC GGCGCAGGGT GCGGTAGGGC GTAGACTGGT GGTCGTTTA

+2 D F C I E L G N K R W Q F N R Q

3601 GGATTTTGC ATCGAGCTGG GTAATAAGCG TTGGCAATT ACCGCCAGT
CCTAAAAACG TAGCTCGACC CATTATTCCG AACCGTTAAA TTGGCGGTCA

+2 S G F L S Q M W I G D K K Q L L T

3651 CAGGCTTTCT TTCACAGATG TGGATTGGCG ATAAAAAACAA ACTGCTGACG
GTCCGAAAGA AAGTGTCTAC ACCTAACCGC TATTTTTGT TGACGACTGC

+2 P L R D Q F T R A P L D N D I G V

3701 CCGCTGCGCG ATCAGTTAC CCGTGCACCG CTGGATAACG ACATTGGCGT
GGCGACGCGC TAGTCAAGTG GGCACGTGGC GACCTATTGC TGTAACCGCA

+2 S E A T R I D P N A W V E R W K

3751 AAGTGAAGCG ACCCGCATTG ACCCTAACGC CTGGGTCGAA CGCTGGAAGG
TTCACTTCGC TGGCGTAAC TGGGATTGCG GACCCAGCTT GCGACCTTCC

+2 A A G H Y Q A E A A L L Q C T A D

3801 CGGCGGGCCA TTACCAAGGCC GAAGCAGCGT TGTTGCAGTG CACGGCAGAT
GCGGCCCGGT AATGGTCCGG CTTCGTCGCA ACAACGTAC GTGCCGTCTA

+2 T L A D A V L I T T A H A W Q H Q

3851 ACACTTGCTG ATGCGGTGCT GATTACGACC GCTCACCGCGT GGCAGCATCA
TGTGAACGAC TACGCCACGA CTAATGCTGG CGAGTGCAC CCGTCGTAGT

+2 G K T L F I S R K T Y R I D G S

3901 GGGGAAAACC TTATTTATCA GCCGGAAAAC CTACCGGATT GATGGTAGTG
CCCCTTTGG AATAAAATAGT CGGCCTTTG GATGCCCTAA CTACCATCAC

+2 G Q M A I T V D V E V A S D T P H

3951 GTCAAATGGC GATTACCGTT GATGTTGAAG TGGCGAGCGA TACACCGCAT
CAGTTACCG CTAATGGCAA CTACAACCTC ACCGCTCGCT ATGTGGCGTA

+2 P A R I G L N C Q L A Q V A E R V

4001 CGGGCGCGGA TTGGCCTGAA CTGCCAGCTG GCGCAGGTAG CAGAGCGGGT
GGCCGCGCCT AACCGGACTT GACGGTCGAC CGCGTCCATC GTCTCGCCCA

+2 N W L G L G P Q E N Y P D R L T

4051 AAACTGGCTC GGATTAGGGC CGCAAGAAA CTATCCGAC CGCCTTACTG
TTTGACCGAG CCTAATCCCG GCGTTCTTT GATAGGGCTG GCGGAATGAC

+2 A A C F D R W D L P L S D M Y T P

4101 CCGCCTGTT TGACCGCTGG GATCTGCCAT TGTCAGACAT GTATACCCCG
GGCGGACAAA ACTGGCGACC CTAGACGGTA ACAGTCTGTA CATATGGGC

+2 Y V F P S E N G L R C G T R E L N

4151 TACGTCTTCC CGAGCGAAAAA CGGTCTGC CGC TGCGGGACGC GCGAATTGAA
ATGCAGAAGG GCTCGCTTTT GCCAGACGCG ACGCCCTGCG CGCTTAACTT

+2 Y G P H Q W R G D F Q F N I S R

4201 TTATGGCCA CACCACTGGC GCGGCAGACTT CCAGTCAAC ATCAGCCGCT
AATACCGGGT GTGGTCACCG CGCCGCTGAA GGTCAAGTTG TAGTCGGCGA

+2 Y S Q Q Q L M E T S H R H L L H A

4251 ACAGTCAACA GCAACTGATG GAAACCAGCC ATCGCCATCT GCTGCACGCG
TGTCAGTTGT CGTTGACTAC CTTTGGTCGG TAGCGGTAGA CGACGTGCGC

+2 E E G T W L N I D G F H M G I G G

4301 GAAGAAGGCA CATGGCTGAA TATCGACGGT TTCCATATGG GGATTGGTGG
CTTCTTCGGT GTACCGACTT ATAGCTGCCA AAGGTATACC CCTAACCAACC

+2 D D S W S P S V S A E F Q L S A

4351 CGACGACTCC TGGAGCCCGT CAGTATCGGC GGAATTCAG CTGAGCGCCG
GCTGCTGAGG ACCTCGGGCA GTCATAGCCG CCTTAAGGTC GACTCGCGGC

+2 G R Y H Y Q L V W C Q K R S D Y K

4401 GTCGCTACCA TTACCAAGTTG GTCTGGTGTG AAAAAAGATC TGACTATAAA
CAGCGATGGT AATGGTCAAC CAGACCACAG TTTTTCTAG ACTGATATT

+2 D E D L D H H H H H R >

4451 GATGAGGACC TCGACCACATCA TCATCATCAT CACCGGTAAT AATAGGTAGA
CTACTCCTGG AGCTGGTAGT AGTAGTAGTA GTGGCCATTA TTATCCATCT

4501 TAAGTGACTG ATTAGATGCA TTGATCCCTC GACCAATTCC GGTTATTTTC
ATTCACTGAC TAATCTACGT AACTAGGGAG CTGGTTAAGG CCAATAAAAG

4551 CACCATATTG CCGTCTTTG GCAATGTGAG GGCCCGGAAA CCTGGCCCTG
GTGGTATAAC GGCAGAAAAC CGTTACACTC CCGGGCCTTT GGACCGGGAC

4601 TCTTCTTGAC GAGCATTCTC AGGGGTCTTT CCCCTCTCGC CAAAGGAATG
AGAAGAACTG CTCGTAAGGA TCCCCAGAAA GGGGAGAGCG GTTTCTTAC

4651 CAAGGTCTGT TGAATGTCGT GAAGGAAGCA GTTCCCTCTGG AAGCTTCTTG
GTTCCAGACCA ACTTACAGCA CTTCCTTCGT CAAGGAGACCC TTCGAAGAAC

4701 AAGACAAACA ACGTCTGTAG CGACCCCTTG CAGGCAGCGG AACCCCCCAC
TTCTGTTGT TGCAAGACATC GCTGGGAAAC GTCCGTCGCC TTGGGGGGTG

4751 CTGGCGACAG GTGCCTCTGC GGCAAAGC CACGTGTATA AGATACACCT
GACCGCTGTC CACGGAGACG CCGGTTTCG GTGCACATAT TCTATGTGGA

4801 GCAAAGCGG CACAACCCA GTGCCACGTT GTGAGTTGGA TAGTTGTGGA
CGTTCCGCC GTGTTGGGGT CACGGTGCAA CACTAACCT ATCAACACCT

4851 AAGAGTCAAA TGGCTCTCCT CAAGCGTATT CAACAAGGGG CTGAAGGATG
TTCTCAGTT ACCGAGAGGA GTTCGCATAA GTTGTCCCC GACTTCCTAC

4901 CCCAGAAGGT ACCCCATTGT ATGGGATCTG ATCTGGGCC TCGGTGCACA
GGGTCTCCA TGGGTAACA TACCCCTAGAC TAGACCCCGG AGCCACGTGT

4951 TGCTTACAT GTGTTAGTC GAGGTTAAAA AACGTCTAGG CCCCCCGAAC
ACGAAATGTA CACAAATCAG CTCCAATTT TTGCAGATCC GGGGGGCTTG

5001 CACGGGGACG TGGTTTCCT TTGAAAAACA CGATGATAAT ACCATGATTG
GTGCCCCCTGC ACCAAAAGGA AACTTTTGT GCTACTATTA TGGTACTAAC

5051 ACAAGATGG ATTGCACGCA GGTTCTCCGG CCGCTTGGGT GGAGAGGCTA
TTGTTCTACC TAACGTGCGT CCAAGAGGCC GGCGAACCCA CCTCTCCGAT

5101 TTCGGCTATG ACTGGGCACA ACAGACAAATC GGCTGCTCTG ATGCCGCCGT
AAGCCGATAC TGACCCGTGT TGTCTGTTAG CCGACGAGAC TACGGCGCA

5151 GTTCCGGCTG TCAGCGCAGG GGCGCCCGGT TCTTTTGTC AAGACCGACC
CAAGGCGAC AGTCGCGTCC CCGCGGCCA AGAAAAACAG TTCTGGCTGG

5201 TGTCCGGTGC CCTGAATGAA CTGCAGGACG AGGCAGCGCG GCTATCGTGG
ACAGGCCACG GGACTTACTT GACGTCTGC TCCGTCGCGC CGATAGCAC

5251 CTGGCCACGA CGGGCGTTCC TTGCGCAGCT GTGCTCGACG TTGTCACTGA
GACCGGTGCT GCCCGCAAGG AACGCGTCGA CACGAGCTGC AACAGTGACT

5301 AGCGGGAAGG GACTGGCTGC TATTGGCGA AGTGGCGGGG CAGGATCTCC
TCGCCCTCC CTGACCGACG ATAACCCGCT TCACGGCCCC GTCCTAGAGG

5351 TGTCACTCTCA CCTTGCTCCT GCCGAGAAAG TATCCATCAT GGCTGATGCA
ACAGTAGAGT GGAACGAGGA CGGCTCTTC ATAGGTAGTA CCGACTACGT

5401 ATGCGGCGGC TGCATACGCT TGATCCGGCT ACCTGCCAT TCGACCACCA
TACGCGCCG ACGTATGCGA ACTAGGCCGA TGGACGGTA AGCTGGTGGT

5451 AGCGAAACAT CGCATCGAGC GAGCACGTAC TCGGATGGAA GCCGGTCTTG
TCGCTTGTA GCGTAGCTCG CTCGTGCATG AGCCTACCTT CGGCCAGAAC

5501 TCGATCAGGA TGATCTGGAC GAAGAGCATC AGGGGCTCGC GCCAGCCGAA
AGCTAGTCCT ACTAGACCTG CTTCTCGTAG TCCCCGAGCG CGGTCGGCTT

5551 CTGTTCGCCA GGCTCAAGGC GCGCATGCC GACGGCGAGG ATCTCGTCGT
GACAAGCGGT CGAGTTCCG CGCGTACGGG CTGCCGCTCC TAGAGCAGCA

5601 GACCCATGGC GATGCCTGCT TGCCGAATAT CATGGTGGAA AATGGCCGCT
CTGGGTACCG CTACGGACGA ACGGCTTATA GTACCACCTT TTACCGGGCA

5651 TTTCTGGATT CATCGACTGT GGCGCGCTGG GTGTGGCGGA CCGCTATCAG
AAAGACCTAA GTAGCTGACA CGGGCCGACC CACACCGCCT GGCGATAGTC

5701 GACATAGCGT TGGCTACCCG TGATATTGCT GAAGAGCTTG GCGGCGAATG
CTGTATCGCA ACCGATGGGC ACTATAACGA CTTCTCGAAC CGCCGCTTAC

5751 GGCTGACCGC TTCTCGTGC TTTACGGTAT CGCCGCTCCC GATTGCGAGC
CCGACTGGCG AAGGAGCACG AAATGCCATA GCGCGAGGG CTAAGCGTCG

5801 GCATCGCCTT CTATCGCCTT CTTGACGAGT TCTCTGAGC GGGACTCTGG
CGTAGCGGAA GATAGCGGAA GAACTGCTCA AGAAGACTCG CCCTGAGACC

5851 GGTCGCATC GATAAAATAA AAGATTTAT TTAGTCTCCA GAAAAAGGGG
CCAAGCGTAG CTATTTTATT TTCTAAAATA AATCAGAGGT CTTTTCCCC

5901 GGAATGAAAG ACCCCACCTG TAGGTTGGC AAGCTAGCTT AAGTAACGCC
CCTTACTTTC TGGGGTGGAC ATCCAAACCG TTCGATCGAA TTCATTGCGG

5951 ATTTTGAAG GCATGGAAA ATACATAACT GAGAATAGAG AAGTTCAGAT
TAAAACGTT CGTACCTTT TATGTATTGA CTCTTATCTC TTCAAGTCTA

6001 CAAGGTCAAG AACAGATGGA ACAGCTGAAT ATGGGCCAAA CAGGATATCT
GTTCCAGTCC TTGTCTACCT TGTCGACTTA TACCCGGTTT GTCCCTATAGA

6051 GTGGTAAGCA GTTCCCTGCC CGGCTCAGGG CCAAGAACAG ATGGAACAGC
CACCATTCTG CAAGGACGGG GCCGAGTCCC GGTTCTTGTC TACCTTGTGCG

6101 TGAATATGGG CCAAACAGGA TATCTGTGGT AAGCAGTTCC TGCCCCGGCT
ACTTATACCC GGTTTGTCT ATAGACACCA TTCGTCAAGG ACGGGGCCGA

6151 CAGGGCAAG AACAGATGGT CCCCAGATGC GGTCCAGCCC TCAGCAGTT
GTCCCCGGTTT TTGTCTACCA GGGGTCTACG CCAGGTCGGG AGTCGTCAAA

6201 CTAGAGAACC ATCAGATGTT TCCAGGGTGC CCCAAGGACC TGAAATGACC
GATCTCTTGG TAGTCTACAA AGGTCCCACG GGGTTCTGG ACTTTACTGG

6251 CTGTGCCCTTA TTTGAACTAA CCAATCAGTT CGCTTCTCGC TTCTGTTGCG
GACACGGAAT AAACCTGATT GGTTAGTCAA GCGAAGAGCG AAGACAAGCG

6301 GCGCTCTGC TCCCCGAGCT CAATAAAAGA GCCCACAACC CCTCACTCGG
CGCGAAGACG AGGGGCTCGA GTTATTTCCT CGGGTGTGTTGG GGAGTGAGCC

6351 GGCGCCAGTC CTCCGATTGA CTGAGTCGCC CGGGTACCCG TGTATCCAAT
CCGCGGTCAG GAGGCTAACT GACTCAGCGG GCCCATGGGC ACATAGGTTA

6401 AAACCCCTTT GCAGTTGCAT CCGACTTGTG GTCTCGCTGT TCCTTGGGAG
TTTGGGAGAA CGTCAACGTA GGCTAACAC CAGAGCGACA AGGAACCCCTC

6451 GGTCTCCTCT GAGTGATTGA CTACCCGTCA GCGGGGGTCT TTCTTCAATG
CCAGAGGAGA CTCACTAACT GATGGGCAGT CGCCCCCAGA AAGTAAGTAC

6501 CAGCATGTAT CAAAATTAAT TTGGTTTTTT TTCTTAAGTA TTTACATTA
GTCGTACATA GTTTAATTA AACCAAAAAA AAGAATTCTA AAATGTAATT

6551 ATGGCCATAG TTGCATTAAT GAATCGGCCA ACGCGCGGGG AGAGGCGGTT
TACCGGTATC AACGTAATT CTTAGCCGGT TCGCGGCCAA TCTCCGCCAA

6601 TCGTATTGG CGCTCTTCG CTTCCCTCGCT CACTGACTCG CTGCGCTCGG
ACGCATAACC GCGAGAAGGC GAAGGAGCGA GTGACTGAGC GACGCGAGCC

6651 TCGTTGGCT GCGCGAGCG GTATCAGCTC ACTCAAAGGC GGTAATACGG
AGCAAGCCGA CGCCGCTCGC CATAGTCGAG TGAGTTCCG CCATTATGCC

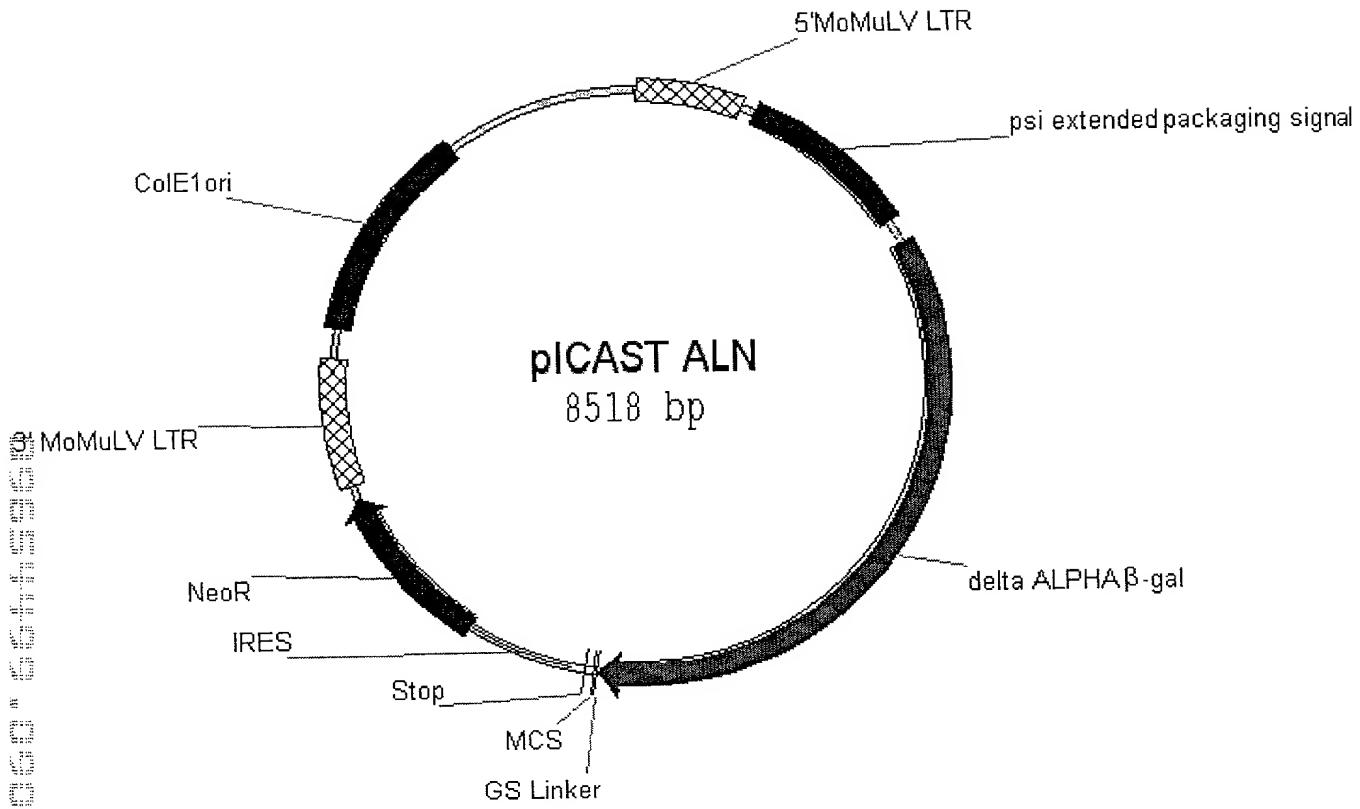


Figure 11A

1 CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
GACGTCGGAC TTATACCCGG TTTGTCTAT AGACACCATT CGTCAAGGAC

51 CCCCCGGCTCA GGGCCAAGAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
GGGGCCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTGT

101 GGATATCTGT GGTAAGCAGT TCCTGCCCG GCTCAGGGCC AAGAACAGAT
CCTATAGACA CCATTCGTCA AGGACGGGGC CGAGTCCCAGG TTCTTGTCTA

151 GGTCCCCAGA TCGGGTCCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
CCAGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGGTAGTCTA

201 GTTTCCAGGG TGCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
CAAAGGTCCC ACGGGGTTCC TGGACTTAC TGGACACGG AATAAACTTG

251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGCT

301 GCTCAATAAA AGAGCCCACA ACCCCCTCACT CGGGGCGCCA GTCCTCCGAT
CGAGTTATTT TCTCGGGTGT TGGGGAGTGA GCGCCCGCGGT CAGGAGGCTA

351 TGACTGAGTC GCCCGGGTAC CCGGTATCC AATAAACCTT CTTGCAGTTG
ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC

401 CATCCGACTT GTGGTCTCGC TGTTCTTGG GAGGGTCTCC TCTGAGTGT
GTAGGCTGAA CACCAGAGCG ACAAGGAACC CTCCCAGAGG AGACTCACTA

451 TGACTACCCG TCAGCGGGGG TCTTCATTT GGGGGCTCGT CCGGGATCGG
ACTGATGGC AGTCGCCCCC AGAAAGTAAA CCCCCGAGCA GGCCCTAGCC

501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
CTCTGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG

551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTAA
TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGACTAAAT

601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG

651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGGCGCAACC CTGGGAGACG
GCACCACCTT GACTGCTCAA GACTTGTGGG CCGGCCTGG GACCCTCTGC

701 TCCCAGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
AGGGTCCCTG AAACCCCCGG CAAAAACACC GGGCTGGACT CCTTCCCTCA

751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGTTCTGGT AGGAGACGAG
GCTACACCTT AGGCTGGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC

801 AACCTAAAAC AGTTCCCGCC TCCGTCTGAA TTTTGCTTT CGGTTGGAA
TTGGATTGG TCAAGGGCGG AGGCAGACTT AAAAACGAAA GCCAAACCTT

851 CCGAAGCCGC GCGTCTTGTC TGCTGCAGCA TCGTTCTGTG TTGTCTGT
GGCTTCGGCG CGCAGAACAG ACGACGTGTC AGCAAGACAC AACAGAGACA

901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACAC
GACTGACACA AAGACATAAA CAGACTTTA ATCCCGGTCT GACAATGGTG

FIGURE 11B

951 TCCCTTAAGT TTGACCTTAG GTAACTGGAA AGATGTCGAG CGGCTCGCTC
AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG

1001 ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
TGTTGGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA

1051 GCAGAACGGC CAACCTTAA CGTCGGATGG CCGCGAGACG GCACCTTAA
CGTCTTACCG GTTGGAAATT GCAGCCTACC GGCGCTCTGC CGTGGAAATT

1101 CCGAGACCTC ATCACCCAGG TTAAGATCAA GGCTTTCA CCTGGCCCGC
GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG

1151 ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCCT TCGGAACCGA

1201 TTTGACCCCC CTCCCTGGGT CAAGCCCTT GTACACCCCTA AGCCTCCGCC
AAACTGGGGG GAGGGACCCA GTTCGGAAA CATGTGGGAT TCGGAGGGCG

1251 TCCTCTTCCT CCATCCGCC CGTCTCTCCC CCTTGAACCT CCTCGTTCGA
AGGAGAAGGA GGTAGCGGG GCAGAGAGGG GGAACTTGGA GGAGCAAGCT

1301 CCCCCGCTCG ATCCTCCCT TATCCAGCCC TCACTCCTTC TCTAGGCCTC
GGGGCGGAGC TAGGAGGGAA ATAGTCGGG AGTGAGGAAG AGATCCCGGG

1351 GGCCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAACACCA
CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCCTA AGCTTGTGGT

1401 TGCACCATCA TCATCATCAC GTCGACTATA AAGATGAGGA CCTCGAGATG
ACGTGGTAGT AGTAGTAGTG CAGCTGATAT TTCTACTCCT GGAGCTCTAC

1451 GGC GTGATTA CGGATTCACT GGCGTCGTG GCCCGCACCG ATCGCCCTTC
CCGCACTAAT GCCTAAGTGA CGGCAGCAC CGGGCGTGGC TAGCGGAAG

1501 CCAACAGTTA CGCAGCCTGA ATGGCGAATG GGCGTTGCC TGGTTTCCGG
GGTTGTCAAT CGT CGGACT TACCGCTTAC CGCGAAACGG ACCAAAGGCC

1551 CACCAAGC GGTGCCGGAA AGCTGGCTGG AGTGCAGATCT TCCTGAGGCC
GTGGTCTTCG CCACGGCCTT TCGACCGACC TCA CGCTAGA AGGACTCCGG

1601 GATACTGTG TCGTCCCCCTC AAACTGGCAG ATGCACGGTT ACGATGCGCC
CTATGACAGC AGCAGGGGAG TTTGACCGTC TACGTGCCAA TGCTACCGGG

1651 CATCTACACC AACGTGACCT ATCCCATTAC GGTCAATCCG CCGTTGTT
GTAGATGTGG TTGCACTGGA TAGGGTAATG CCAGTTAGGC GGAAACAAG

1701 CCACGGAGAA TCCGACGGGT TGTTACTCGC TCACATTTAA TGTTGATGAA
GGTGCCTCTT AGGCTGCCA ACAATGAGCG AGTGTAAATT ACAACTACTT

1751 AGCTGGCTAC AGGAAGGCCA GACGCGAATT ATTTTGATG GCGTTAACTC
TCGACCGATG TCCCTCCGGT CTGCGCTTAA TAAAAACTAC CGCAATTGAG

1801 GGC GTTTCAT CTGTGGTGCA ACGGCGCTG GGTGGTTAC GGCGAGGACA
CCGCAAAGTA GACACCACGT TGCCCGCGAC CCAGCCAATG CGGGCCTGT

1851 GTCGTTGCC GTCTGAATT GACCTGAGCG CATTGGTACG CGCCGGAGAA
CAGCAAACGG CAGACTAAA CTGGACTCGC GTAAAAATGC CGGGCCTTT

1901 AACCGCCTCG CGGTGATGGT GCTGCGCTGG AGTGACGGCA GTTATCTGGA
TTGGCGGAGC GCCACTACCA CGACGCGACC TCACTGCCGT CAATAGACCT

1951 AGATCAGGAT ATGTGGCGGA TGAGCGGCAT TTTCCGTGAC GTCTCGTTGC
TCTAGTCCTA TACACCGCCT ACTCGCCGTAA AAAGGCACGT CAGAGCAACG

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2201 CGTGGTGGTT ATGCCGATCG CGTCACACTA CGTCTGAACG TCGAAAACCC
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2251 GAAAATGTGG AGCGCCGAAA TCCCGAATCT CTATCGTGC G TGGGTTGAAAC
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2451 AGGTCACTGGA TGAGCAGACG ATGGTGCAGG ATATCCTGCT GATGAAGCAG
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2901 TGCCGAAATG GTCCCATAAA AAATGGCTTT CGCTACCTGG AGAGACGCGC
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3201 CAGCGCTGAC GGAAGCAAAA CACCAGCAGC AGTTTTCCA GTTCCGTTA
GTCGCGACTG CCTTCGTTT GTGGTCGTG TCAAAAAGGT CAAGGCAAAT

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CGCCACTTCA CGGAGACCTA CAGCGAGGTG TTCCATTGT CAACTAACTT

3401 CTGCTGAAC TACCGCAGCC GGAGAGCGCC GGGCAACTCT GGCTCACAGT
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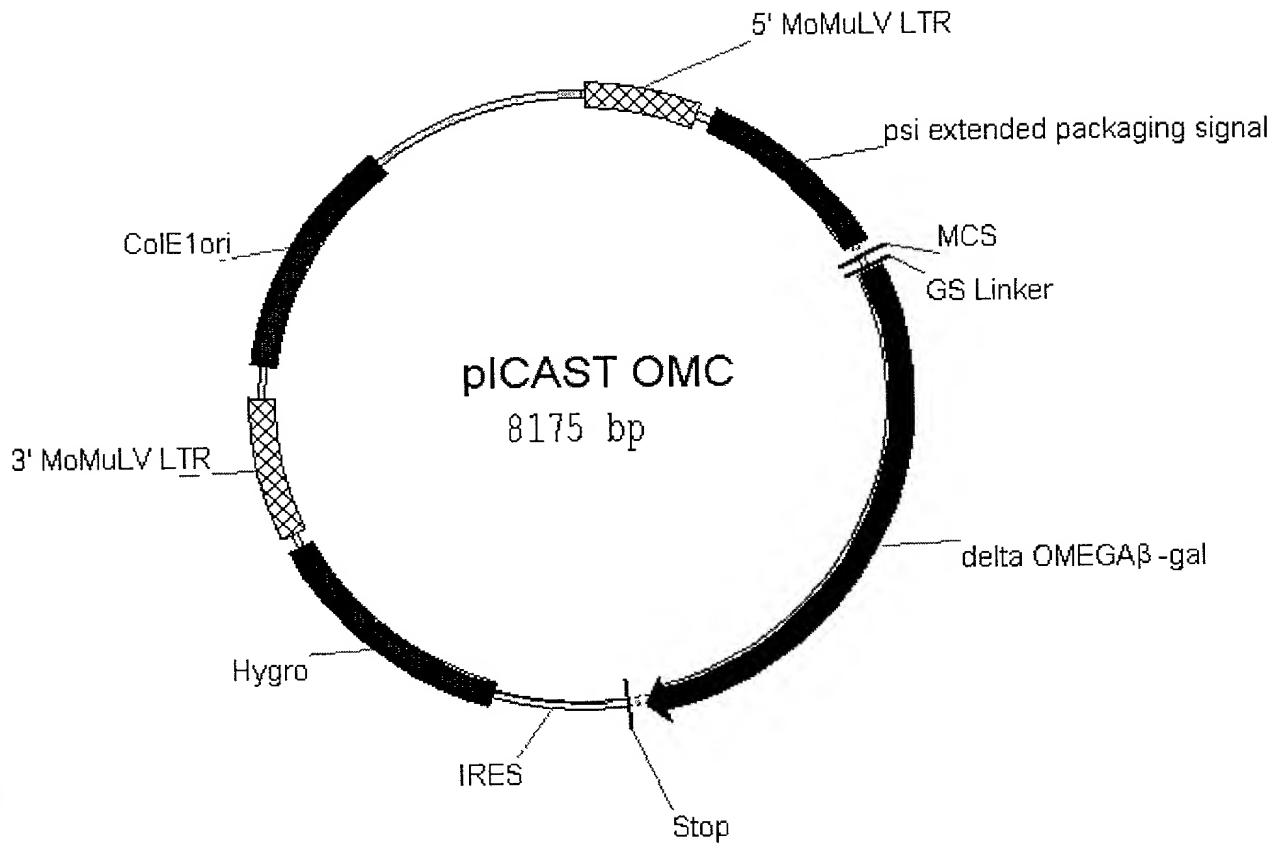


Figure 12A

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CTCTGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG

551 AGCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTAA
TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGACTAAAAT

601 TGCCTCGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCTGG

651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGCGCAACC CTGGGAGACG
GCACCACCTT GACTGCTCAA GACTTGTGGG CGCGCGTTGG GACCCTCTGC

701 TCCCAGGGAC TTTGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
AGGGTCCCTG AAACCCCCGG CAAAAACACC GGGCTGGACT CCTCCCTCA

751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
GCTACACCTT AGGCTGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC

801 AACCTAAAAC AGTCCCCGCC TCCGCTGAA TTTTTGCTTT CGGTTGGAA
TTGGATTTG TCAAGGGCGG AGGCAGACTT AAAAACGAAA GCCAAACCTT

851 CCGAAGCCGC GCGTCTTGTG TGCTGCAGCA TCGTTCTGTG TTGTCTCTGT
GGCTCGGGCG CGCAGAACAG ACGACGTGAGT AGCAAGACAC AACAGAGACA

901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACCAC
GACTGACACA AAGACATAAA CAGACTTTA ATCCCGGTCT GACAATGGTG

FIGURE 12B

951 TCCCTTAAGT TTGACCTTAG GTAACTGGAA AGATGTCGAG CGGCTCGCTC
AGGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG

1001 ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
TGTTGGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGAGA

1051 GCAGAACGGC CAACCTTAA CGTCGGATGG CCGCGAGACG GCACCTTAA
CGTCTTACCG GTTGGAAATT GCAGCCTACC GGCGCTCTGC CGTGGAAATT

1101 CCGAGACCTC ATCACCCAGG TTAAGATCAA GGTCTTTCA CCTGGCCCGC
GGCTCTGGAG TAGTGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG

1151 ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTTGGCT
TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCCT TCGGAACCGA

1201 TTTGACCCCC CTCCCTGGGT CAAGCCCTT GTACACCCTA AGCCTCCGCC
AAACTGGGG GAGGGACCCA GTTCGGAAA CATGTGGAT TCGGAGGCCG

1251 TCCTCTTCCCT CCATCCGCC CGTCTCTCCC CCTGGAACCT CCTCGTTCGA
AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTTGGGA GGAGCAAGCT

1301 CCCCGCTCTG ATCCCTCCCTT TATCCAGCCC TCACTCCCTC TCTAGGCC
GGGGCGGAGC TAGGAGGGAA ATAGTCGGG AGTGAGGAAG AGATCCCGG

1351 GGCCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAATCAGG
CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTAGTCC

1401 CCTTGGCGCG CGGGATCCTT AATTAAGCGC AATTGGGAGG TGGCGGTAGC
GGAACCGCGC GGCCTAGGAA TTAATTCGCG TTAACCCCTCC ACCGCCATCG

1451 CTCGAGATGG GCGTGATTAC GGATTCACTG GCCGTCGTTT TACAACGTG
GAGCTCTACC CGCACTAATG CCTAAGTGAC CGGCAGCAAA ATGTTGCAGC

1501 TGACTGGAA AACCCCTGGCG TTACCCAAT TAATCGCTT GCAGCACATC
ACTGACCCCTT TTGGGACCGC AATGGGTGA ATTAGCGGAA CGTCGTGTAG

1551 CCCCTTCGC CAGCTGGCGT AATAGCGAAG AGGCCGCAC CGATGCCCT
GGGGAAAGCG GTCGACCGCA TTATCGCTTC TCCGGCGTG GCTAGCGGG

1601 TCCCCAACAGT TACGCAGCCT GAATGGCGAA TGGCGCTTTG CCTGGTTCC
AGGGTTGTCA ATGCGTCGGA CTTACCGCTT ACCCGCAAAC GGACCAAAGG

1651 GGCACCCAGAA GCGGTGCCGG AAAGCTGGCT GGAGTGCAGT CTTCTGAGG
CCGTGGTCTT CGCCACGGCC TTTCGACCGA CCTCACGCTA GAAGGACTCC

1701 CCGATACTGT CGTCGTCCCC TCAAACCTGGC AGATGCACGG TTACGATGCG
GGCTATGACA GCAGCAGGGG AGTTTGACCG TCTACGTGCC AATGCTACGC

1751 CCCATCTACA CCAACGTGAC CTATCCCATT ACGGTCAATC CGCCGTTGT
GGGTAGATGT GGTTGCAGT GATAGGGTAA TGCCAGTTAG GCGGCAAACA

1801 TCCCACGGAG AATCCGACGG GTTGTACTC GCTCACATT AATGTTGATG
AGGGTGCCTC TTAGGCTGCC CAACAATGAG CGAGTGTAAA TTACAACATAC

1851 AAAGCTGGCT ACAGGAAGGC CAGACCGCAGA TTATTTTGA TGGCGTTAAC
TTTCGACCGA TGTCCCTTCG GTCTGCGCTT AATAAAAACCT ACCGCAATTG

1901 TCGGCGTTTC ATCTGTGGTG CAACGGGCAGC TGGGTCGGTT ACGGCCAGGA
AGCCGAAAG TAGACACCAC GTTGCCTCGC ACCCAGCCAA TGCCGGTCCT

1951 CAGTCGTTTG CCGTCTGAAT TTGACCTGAG CGCATTCTTA CGCGCCGGAG
GTCAGCAAAC GGCAGACTTA AACTGGACTC GCGTAAAAAT GCGCGGCCTC

2001 AAAACCGCCT CGCGGTGATG GTGCTGCCGT GGAGTGACGG CAGTTATCTG
TTTTGGCGGA CGCCCACTAC CACGACCGCA CCTCACTGCC GTCAATAGAC

2051 GAAGATCAGG ATATGTGGCG GATGAGCGGC ATTTCCTCGT ACgtCTCGTT
CTTCTAGTCC TATAACCCGC CTACTCGCCG TAAAAGGCAC TGCAAGAGCAA

2101 GCTGCATAAA CCGACTACAC AAATCAGCGA TTTCCATGTT GCCACTCGCT
CGACGTATTT GGCTGATGTG TTTAGTCGCT AAAGGTACAA CGGTGAGCGA

2151 TTAATGATGA TTTCAGCCGC GCTGTACTGG AGGCTGAAGT TCAGATGTGC
AATTACTACT AAAGTCGGCG CGACATGACC TCCGACTTCA AGTCTACACG

2201 GGCAGGTTGC GTGACTACCT ACGGGTAACA GTTCTTTAT GGCAGGGTGA
CCGCTCAACG CACTGATGGA TGCCCATTGT CAAAGAAATA CCGTCCCAC

2251 AACGCAGGTC GCCAGCGGC CCGCGCCTT CGGCAGGTGAA ATTATCGATG
TTGCCTCCAG CGGTGCGCCGT CGCGCGAAA GCCGCCACTT TAATAGCTAC

2301 AGCGTGGTGG TTATGCCAT CGCGTCACAC TACGTCTGAA CGTCGAAAAC
TCGCACCAACC AATACGGCTA CGCGAGTG TG ATGCAGACTT GCAGCTTTG

2351 CCGAAACTGT GGAGCGCCGA AATCCCGAAT CTCTATCGT CGGTGGTTGA
GGCTTGACA CCTCGCGGCT TTAGGGCTTA GAGATAGCAC GCCACCAACT

2401 ACTGCACACC GCCGACGGCA CGCTGATTGA AGCAGAAGCC TGCGATGTCG
TGACGTGTGG CGGCTGCCGT GCGACTAACT TCGTCTCGG ACGCTACAGC

2451 GTTCCCGCGA GGTGCGGATT GAAAATGGTC TGCTGCTGCT GAACGGCAAG
CAAAGGCCTA CCACGCCTAA CTTTACCAAG ACCACGACGA CTTGCCGTT

2501 CCGTTGCTGA TTCGAGGCCT TAACCGTCAC GAGCATCATC CTCTGCATGG
GGCAACGACT AAGCTCCGCA ATTGGCAGTG CTCGTAGTAG GAGACGTACC

2551 TCAGGGTCATG GATGAGCAGA CGATGGTGCA GGATATCCTG CTGATGAAGC
AGTCCAGTAC CTACTCGTCT GCTACCACGT CCTATAGGAC GACTACTCG

2601 AGAACAACTT TAACGCCGTG CGCTGTTCGC ATTATCCGAA CCATCCGCTG
TCTTGTGAA ATTGCGGCAC GCGACAAGCG TAATAGGCTT GGTAGGGCAG

2651 TGGTACACGC TGTGCGACCG CTACGGCCTG TATGTGGTGG ATGAAGCCAA
ACCATGTGCG ACACGCTGGC GATGCCGGAC ATACACCACCA TACTTCGGTT

2701 TATTGAAACC CACGGCATGG TGCCAATGAA TCGTCTGACC GATGATCCGC
ATAACCTTGG GTGCCGTACC ACGGTTACTT AGCAGACTGG CTACTAGGCG

2751 GCTGGCTACC GGCAGATGAGC GAACGCGTAA CGCGAATGGT GCAGCGCGAT
CGACCGATGG CGCCTACTCG CTTGCCGATT GCGCTTACCA CGTCGCGCTA

2801 CGTAATCACC CGAGTGTGAT CATCTGGTCG CTGGGGAATG AATCAGGCCA
GCATTAGTGG GCTCACACTA GTAGACCAGC GACCCCTTAC TTAGTCCGGT

2851 CGGCCTAAT CACGACGCGC TGTATCGCTG GATCAAATCT GTCGATCCTT
GCCGCGATTA GTGCTGCGCG ACATAGCGAC CTAGTTAGA CAGCTAGGAA

2901 CCCGCCGGT GCAGTATGAA GGCGCCGGAG CCGACACCCAC GGCCACCGAT
GGGCAGGCCA CGTCATACTT CCGCCGCCCTC GGCTGTGGTG CCGGTGGCTA

2951 ATTATTTGCC CGATGTACGC GCGCGTGGAT GAAGACCCAGC CCTTCCCCGC
TAATAAACGG GCTACATGCG CGCGCACCTA CTTCTGGTCG GGAAGGGCCG

3001 TGTGCCGAAA TGGTCCATCA AAAAATGGCT TTCGCTACCT GGAGAGACGC
ACACGGCTTT ACCAGGTAGT TTTTACCGA AAGCGATGGA CCTCTCTGCG

3051 GCCCGCTGAT CCTTTGCGAA TACGCCACG CGATGGTAA CAGTCTTGGC
CGGGCGACTA GGAAACGCTT ATGCGGGTGC GCTACCCATT GTCAGAACCG

3101 GGTTTCGCTA AATACTGGCA GGCGTTTCGT CAGTATCCCC GTTTACAGGG
CCAAAGCGAT TTATGACCGT CCGCAAAGCA GTCATAGGGG CAAATGTCCC

3151 CGGCTCGTC TGGGACTGGG TGGATCAGTC GCTGATTAAA TATGATGAAA
GCCGAAGCAG ACCCTGACCC ACCTAGTCAG CGACTAATT ATACTACTTT

3201 ACGGCAACCCC GTGGTCGGCT TACGGCGGTG ATTTTGGCGA TACGCCGAAC
TGCGTTGGG CACCAGCCGA ATGCCGCCAC TAAAACCGCT ATGCGGCTTG

3251 GATGCCAGT TCTGTATGAA CGGTCTGGTC TTTGCCGACC GCACGCCGA
CTAGCGGTCA AGACATACTT GCCAGACCAAG AAACGGCTGG CGTGCAGCGT

3301 TCCAGCGCTG ACGGAAGCAA AACACCAGCA GCAGTTTTTC CAGTTCCGTT
AGGTCCGAC TGCTTCGTT TTGTGGTCGT CGTCAAGGCAA

3351 TATCCGGGCA AACCATCGAA GTGACCAGCG AATACCTGTT CCGTCATAGC
ATAGGCCCCTT TTGGTAGCTT CACTGGTCGC TTATGGACAA GGCAAGTATCG

3401 GATAACGAGC TCCTGCCTG GATGGTGGCG CTGGATGGTA AGCCGCTGGC
CTATTGCTCG AGGACGTGAC CTACCACCGC GACCTACCAT TCGCGACCG

3451 AAGCGGTGAA GTGCCCTCTGG ATGTCGCTCC ACAAGGTAAA CAGTTGATTG
TTCGCCACTT CACGGAGACC TACAGCGAGG TGTTCATT GTCAACTAAC

3501 AACTGCCCTGA ACTACCGCAG CGGGAGAGCG CCAGGGCAACT CTGGCTCACA
TTGACGGACT TGATGGCGTC GGCCTCTCGC GGCGCGTTGA GACCGAGTGT

3551 GTACCGTAG TGCAACCGAA CGCGACCGCA TGGTCAGAAG CCAGGGCACAT
CATGCGCATC ACGTTGGCTT GCGCTGGCGT ACCAGTCTTC GGCGCGTGT

3601 CAGCGCTGG CAGCACTGGC GTCTGGCGGA AAACCTCAGT GTGACGCTCC
GTCGCGGACC GTCGTCACCG CAGACCGCCT TTTGGAGTC CACTGCGAGG

3651 CCGCCGCGTC CCACGCCATC CCGCATCTGA CCACCCAGCGA AATGGATTT
GGCGCGCGAG GGTGCGGTAG GGCCTAGACT GGTGGTCGCT TTACCTAAAA

3701 TGCATCGAGC TGGGTAATAA GCGTTGGCAA TTTAACCGCC AGTCAGGCTT
ACGTAGCTCG ACCCATTATT CGCAACCGTT AAATTGGCGG TCAGTCCGAA

3751 TCTTTCACAG ATGTGGATTG GCGATAAAAA ACAACTGCTG ACGCCGCTGC
AGAAAGTGTG TACACCTAAC CGCTATTTT TGTTGACGAC TGCAGGCGACG

3801 GCGATCAGTT CACCCGTGTC GATAGATCTG AACAGAAACT CATTTCGAA
CGCTAGTCAA GTGGGCACAG CTATCTAGAC TTGTCTTGA GTAAAGGCTT

3851 GAAGACCTAG TCGACCACATCA TCATCATCAT CACCGGTAAT AATAGGTAGA
CTTCTGGATC AGCTGGTAGT AGTAGTAGTA GTGGCCATTA TTATCCATCT

3901 TAAGTGACTG ATTAGATGCA TTTCGACTAG ATCCCTCGAC CAATTCCGGT
ATTCACTGAC TAATCTACGT AAAGCTGATC TAGGGAGCTG GTTAAGGCCA

3951 TATTTTCCAC CATATTGCCG TCTTTGGCA ATGTGAGGGC CCGGAAACCT
ATAAAAGGTG GTATAACGGC AGAAAACCGT TACACTCCCG GCCCTTTGGA

4001 GGCCCTGTCT TCTTGACGAG CATTCCCTAGG GGTCTTTCCC CTCTGCCAA
CCGGGACAGA AGAACTGCTC GTAAGGATCC CCAGAAAGGG GAGAGCGGTT

4051 AGGAATGCAA GGTCTGTTGA ATGTCGTGAA GGAAGCAGTT CCTCTGGAAG
TCCTTACGTT CCAGACAAC TACAGCACTT CCTTCGTCAA GGAGACCTTC

4101 CTTCTGAAG ACAAAACAACG TCTGTAGCGA CCCTTGCAG GCAGCGGAAC
GAAGAACTTC TGTTTGTGAG AGACATCGCT GGGAAACGTC CGTCGCCTG

4151 CCCCCCACCTG GCGACAGGTG CCTCTGCGGC CAAAAGCCAC GTGTATAAGA
GGGGGTGGAC CGCTGTCCAC GGAGACGCCG GTTTCGGTG CACATATTCT

4201 TACACCTGCA AAGGCGGCAC AACCCCCAGTG CCACGTTGTG AGTTGGATAG
ATGTGGACGT TTCCGCCGTG TTGGGGTCAC GGTGCAACAC TCAACCTATC

4251 TTGTGGAAAG AGTCAAATGG CTCTCCTCAA GCGTATTCAA CAAGGGGCTG
AACACCTTTC TCAGTTTACG GAGAGGAGTT CGCATAAGTT GTTCCCCGAC

4301 AAGGATGCCA AGAAGGTACC CCATTGTATG GGATCTGATC TGGGGCCTCG
TTCCTACGGG TCTTCATGG GGTAACATAC CCTAGACTAG ACCCCGGAGC

4351 GTGCACATGC TTTACATGTG TTTAGTCGAG GTAAAAAAAC GTCTAGGCC
CACGTGTACG AAATGTACAC AAATCAGCTC CAATTTTTG CAGATCCGGG

4401 CCCGAACAC GGGGACGTGG TTTCCCTTG AAAAACACGA TGATAATACC
GGGCTTGGTG CCCCTGCACC AAAAGGAAAC TTTTTGTGCT ACTATTATGG

4451 ATGAAAAAGC CTGAACTCAC CGCGACGTCT GTCGAGAAGT TTCTGATCGA
TACTTTTCG GACTTGAGTG GCGCTGCAGA CAGCTCTCA AAGACTAGCT

4501 AAAGTTCGAC AGCGTCTCCG ACCTGATGCA GCTCTCGGAG GGCAGAGAAT
TTTCAAGCTG TCGCAGAGGC TGGACTACGT CGAGAGCCTC CCGCTCTTA

4551 CTCGTGCTTT CAGCTTCGAT GTAGGGAGGGC GTGGATATGT CCTGGGGTA
GAGCACGAAA GTCGAAGCTA CATCCTCCCG CACCTATACA GGACGCCAT

4601 AATAGCTGCG CCGATGGTT CTACAAAGAT CGTTATGTTT ATCGGCACCTT
TTATCGACGC GGCTACCAAA GATGTTCTA GCAATACAAA TAGCCGTGAA

4651 TGCATCGGCC GCGCTCCCGA TTCCGGAAGT GCTTGACATT GGGGAATTAA
ACGTAGCCGG CGCGAGGGCT AAGGCCTCA CGAAGCTGTAA CCCCTAAAT

4701 GCGAGAGCCT GACCTATTGC ATCTCCGCC GTGCACAGGG TGTCACGTTG
CGCTCTCGGA CTGGATAACG TAGAGGGCGG CACGTGTCCC ACAGTGCAAC

4751 CAAGACCTGC CTGAAACCGA ACTGCCGCT GTTCTGCAGC CGGTCGCGGA
GTTCTGGACG GACTTTGGCT TGACGGCGA CAAGACGTG CCGAGCGCCT

4801 GGCCATGGAT GCGATCGCT CGGCCGATCT TAGCCAGACG AGCGGGTTCG
CCGGTACCTA CGCTAGCGAC GCCGGCTAGA ATCGGTCTGC TCGCCCAAGC

4851 GCCCATTGG ACCGCAAGGA ATCGGTCAT ACACATACATG GCGTGATTTC
CGGGTAAGCC TGGCGTTCCCT TAGCCAGTTA TGTGATGTAC CGCACTAAAG

4901 ATATGCGCGA TTGCTGATCC CCATGTGTAT CACTGGCAAA CTGTGATGGA
TATAACGCGCT AACGACTAGG GGTACACATA GTGACCGTTT GACACTACCT

4951 CGACACCGTC AGTGCCTCCG TCGCGCAGGC TCTCGATGAG CTGATGCTTT
GCTGTGGCAG TCACGCAGGC AGCGCGTCCG AGAGCTACTC GACTACGAAA

5001 GGGCCGAGGA CTGCCCCGAA GTCCGGCACC TCGTGCACGC GGATTTGGC
CCC GGCTCCT GACGGGGCTT CAGGCCGTGG AGCACGTGCG CCTAAAGCCG

5051 TCCAACAATG TCCTGACGGA CAATGGCCGC ATAACAGCGG TCATTGACTG
AGGTTGTTAC AGGACTGCCT GTTACCGCGG TATTGTCGCC AGTAACGTGAC

5101 GAGCGAGGCG ATGTTGGGG ATTCCAATA CGAGGTCGCC AACATCTTCT
CTCGCTCCGC TACAAGCCCC TAAGGGTTAT GCTCCAGCGG TTGTAGAAGA

5151 TCTGGAGGCC GTGGTTGGCT TGTATGGAGC AGCAGACGCG CTACTTCGAG
AGACCTCCGG CACCAACCGA ACATACCTCG TCGTCTGCGC GATGAAGCTC

5201 CGGAGGCATC CGGAGCTTGC AGGATCGCCG CGGCTCCGGG CGTATATGCT
GCCTCCGTAG GCCTCGAACG TCCTAGCGGC GCCGAGGCC GCATATACGA

5251 CCGCATTGGT CTTGACCAAC TCTATCAGAG CTTGGTTGAC GGCAATTTCG
GGCGTAACCA GAACTGGTTG AGATAGTCTC GAACCAACTG CCGTTAAAGC

5301 ATGATGCAGC TTGGGGCGAG GGTCGATGCG ACGCAATCGT CCGATCCGA
TACTACGTG AACCCCGCTC CCAGCTACGC TCGTTAGCA GGCTAGGCCT

5351 GCCGGGACTG TCGGGCGTAC ACAAAATCGCC CGCAGAAAGCG CGGCCGTCTG
CGGCCCTGAC AGCCCGCATG TGTTTACCGG GCGTCTTCGC GCCGGCAGAC

5401 GACCGATGGC TGTGTAGAAC TACTCGCCGA TAGTGGAAAC CGACGCCCCA
CTGGCTACCG ACACATCTTC ATGAGCGGCT ATCACCTTG GCTGCGGGT

5451 GCACTCGTCC GAGGGCAAAG GAATAGAGTA GATGCCGACC GGGATCTATC
CGTGAGCAGG CTCCCCTTTC CTTATCTCAT CTACGGCTGG CCCTAGATAG

5501 GATAAAATAA AAGATTTAT TTAGTCTCCA GAAAAAGGGG GGAATGAAAG
CTATTTTATT TTCTAAAATA AATCAGAGGT CTTTTTCCCC CCTTACTTTC

5551 ACCCCACCTG TAGGTTGGC AAGCTAGCTT AAGTAACGCC ATTTGCAAG
TGGGGTGGAC ATCCAAACCG TTCGATCGAA TTCATTGCGG TAAAACGTTC

5601 GCATGGAAAA ATACATAACT GAGAATAGAG AAGTTCAGAT CAAGGTCAAG
CGTACCTTT TATGTATTGA CTCTTATCTC TTCAAGTCTA GTTCCAGTCC

5651 AACAGATGGA ACAGCTGAAT ATGGGCCAAA CAGGATATCT GTGGTAAGCA
TTGCTCACCT TGTCGACTTA TACCCGGTTT GTCCTATAGA CACCATTGCT

5701 GTTCCTGCC CGGCTCAGGG CCAAGAACAG ATGGAACAGC TGAATATGGG
CAAGGACGGG GCCGAGTCCC GGTTCTGTC TACCTTGTG ACTTATAACCC

5751 CCAAACAGGA TATCTGTGGT AAGCAGTTCC TGCCCCGGCT CAGGGCCAAG
GGTTTGCCT ATAGACACCA TTCGTCAAGG ACGGGGCCGA GTCCCGGTT

5801 AACAGATGGT CCCCAGATGC GGTCCAGCCC TCAGCAGTTT CTAGAGAAC
TTGTCTACCA GGGGTCTACG CCAGGTCGGG AGTCGTCAA GATCTTGTG

5851 ATCAGATGTT TCCAGGGTGC CCCAAGGACC TGAAATGACC CTGTGCCCTA
TAGTCTACAA AGGTCCCACG GGGTTCTGG ACTTTACTGG GACACGGAAT

5901 TTTGAACTAA CCAATCAGTT CGCTTCTCGC TTCTGTTCGC GCGCTTCTGC
AAACTTGATT GGTTAGTCAA GCGAAGAGCG AAGACAAGCG CGCGAAGACG

5951 TCCCCGAGCT CAATAAAAGA GCCCACAAACC CCTCACTCGG GGCGCCAGTC
AGGGGCTCGA GTTATTTCT CGGGTGTGAGCC CGCGGGTCA

6001 CTCCGATTGA CTGAGTCGCC CGGGTACCCG TGATCCAAT AAACCCCTTT
GAGGCTAACT GACTCAGCGG GCCCATGGGC ACATAGGTTA TTTGGGAGAA

6051 GCAGTTGCAT CCGACTTGTG GTCTCGCTGT TCCTGGGAG GGTCTCCCT
CGTCAACGTA GGCTGAACAC CAGAGCGACA AGGAACCCCTC CCAGAGGAGA

6101 GAGTGATTGA CTACCCGTCA GCGGGGGTCT TTCATTATCG CAGCATGTAT
CTCACTAACT GATGGGCAGT CGCCCCCAGA AAGTAAGTAC GTCGTACATA

6151 CAAAATTAAT TTGGTTTTTT TTCTTAAGTA TTACATTAA ATGGCCATAG
GTTTAATTA AACCAAAAAA AAGAATTAT AAATGTAATT TACCGGTATC

6201 TTGCTTAAT GAATCGGCCA ACGCGGGGG AGAGGCGGTT TGCGTATTGG
AACGTAATTA CTTAGCCGGT TGCGGCCAA ACACATAACC

6251 CGCTCTTCG CTTCTCGCT CACTGACTCG CTGCGCTCG TCGTTGGCT
GCGAGAAGGC GAAGGAGCGA GTGACTGAGC GACGCGAGCC AGCAAGCCGA

6301 GCGCGAGCG GTATCAGCTC ACTCAAAGGC GTAAATACGG TTATCCACAG
CGCCGCTCGC CATAGTCGAG TGAGTTCCG CCATTATGCC AATAGGTGTC

6351 AATCAGGGGA TAACGCAGGA AAGAACATGT GAGCAAAGG CCAGCAAAG
TTAGTCCCT ATTGCGTCCT TTCTGTACA CTCGTTTCC GGTGTTTTC

6401 GCCAGGAACC GTAAAAGGC CGCGTTGCTG GCGTTTTCC ATAGGCTCCG
CGGTCTTGG CATTTCGCG GCGCAACGAC CGCAAAAGG TATCCGAGGC

6451 CCCCCCTGAC GAGCATACA AAAATCGACG CTCAAGTCAG AGGTGGCGAA
GGGGGGACTG CTCGTAGTGT TTTAGCTGC GAGTTCACTC TCCACCGCTT

6501 ACCCGACAGG ACTATAAAGA TACCAGGCGT TTCCCCCTGG AAGCTCCCTC
TGGGCTGTCC TGATATTCT ATGGTCCGCA AAGGGGGACC TTCGAGGGAG

6551 GTGCGCTCTC CTGTTCCGAC CCTGCCGCTT ACCGGATACC TGTCCGCCTT
CACGCGAGAG GACAAGGCTG GGACGGCGAA TGGCCTATGG ACAGGCGGAA

6601 TCTCCCTTCG GGAAGCGTGG CGCTTTCTCA TAGCTCACGC TGTAGGTATC
AGAGGGAAGC CCTTCGCAAC GCGAAAGAGT ATCGAGTGCACATCCATAG

6651 TCAGTTCGGT GTAGGTCGTT CGCTCCAAGC TGGGCTGTGT GCACGAACCC
AGTCAAGCCA CATCCAGCAA GCGAGGTTCG ACCCGACACA CGTGCTGGG

6701 CCCGTTCAGC CCGACCGCTG CGCCTTATCC GGTAACTATC GTCTTGAGTC
GGGCAAGTCG GGCTGGCGAC GCGGAATAGG CCATTGATAG CAGAACTCAG

6751 CAACCCGGTA AGACACCGACT TATGCCACT GCCAGCAGCC ACTGGTAACA
GTTGGGCCAT TCTGTGCTGA ATAGCGGTGA CCGTCGTCGG TGACCATTGT

6801 GGATTAGCAG AGCGAGGTAT GTAGGGCGGTG CTACAGAGTT CTTGAAGTGG
CCTAATCGTC TCGCTCCATA CATCCGCCAC GATGTCTCAA GAACTTCACC

6851 TGGCCTAACT ACGGCTACAC TAGAAGAACAA GTATTTGGTA TCTGCCTCT
ACCGGATTGA TGCCGATGTG ATCTTCTTGT CATAAAACCAT AGACGCGAGA

6901 GCTGAAGCCA GTTACCTTCG GAAAAAGAGT TGGTAGCTCT TGATCCGGCA
CGACTTCGGT CAATGGAAGC CTTTTCTCA ACCATCGAGA ACTAGGCCGT

6951 AACAAACAC CGCTGGTAGC GGTGGTTTT TTGTTGCAA GCAGCAGATT
TTGTTTGGTG GCGACCATCG CCACCAAAAA AACAAACGTT CGTCGTCTAA

7001 ACGCGCAGAA AAAAAGGATC TCAAGAACAGT CCTTTGATCT TTTCTACGGG
TGCGCGTCTT TTTTCCTAG AGTTCTTCTA GGAAACTAGA AAAGATGCC

7051 GTCTGACGCT CAGTGGAACG AAAACTCACG TTAAGGGATT TTGGTCATGA
CAGACTGCGA GTCACCTTGC TTTGAGTGC ATTCCCTAA AACCACTACT

7101 GATTATCAA AAGGATCTTC ACCTAGATCC TTTTAAATTAA AAAATGAAGT
CTAATAGTTT TTCCTAGAAG TGGATCTAGG AAAATTTAA TTTTACTTCA

7151 TTGCGGCCGC AAATCAATCT AAAGTATATA TGAGTAAACT TGGTCTGACA
AACGCCGGCG TTTAGTTAGA TTTCATATAT ACTCATTGAA ACCAGACTGT

7201 GTTACCAATG CTTAATCAGT GAGGCACCTA TCTCAGCGAT CTGTCTATT
CAATGGTTAC GAATTAGTCA CTCCGTGGAT AGAGTCGCTA GACAGATAAA

7251 CGTTCATCCA TAGTTGCCGTG ACTCCCCGTC GTGTAGATAA CTACGATACG
GCAAGTAGGT ATCAACGGAC TGAGGGGCAG CACATCTATT GATGCTATGC

7301 GGAGGGCTTA CCATCTGGCC CCAGTGCTGC AATGATACCG CGAGACCCAC
CCTCCGAAT GGTAGACCGG GGTACAGACG TTACTATGGC GCTCTGGGTG

7351 GCTCACCGGC TCCAGATTAA TCAGCAATAA ACCAGCCAGC CGGAAGGGCC
CGAGTGGCCG AGGTCTAAAT AGTCGTTATT TGGTCGGTGC GCCTTCCCGG

7401 GAGCCAGAA GTGGTCCTGC AACTTTATCC GCCTCCATCC AGTCTATTAA
CTCGCGTCTT CACCAGGACG TTGAAATAGG CGGAGGTAGG TCAGATAATT

7451 TTGTTGCCGG GAAGCTAGAG TAAGTAGTTC GCCAGTTAAAT AGTTTGCAGCA
AACAAACGGCC CTTCGATCTC ATTCAATCAAG CGGTCAATTAA TCAAACCGGT

7501 ACGTTGTTGC CATTGCTACA GGCATCGTGG TGTCACGCTC GTCGTTGGT
TGCAACAAACG GTAACGATGT CCGTAGCACC ACAGTGCAG CAGCAAACCA

7551 ATGGCTTCAT TCAGCTCCGG TTCCCAACGA TCAAGGCAGG TTACATGATC
TACCGAAGTA AGTCGAGGCC AAGGGTTGCT AGTCCGCTC AATGTACTAG

7601 CCCCCATGTTG TGCACAAAAG CGGTTAGCTC CTTCGGTCTT CCGATCGTTG
GGGGTACAAC ACGTTTTTC GCCAATCGAG GAAGCCAGGA GGCTAGCAAC

7651 TCAGAAGTAA GTTGGCCGCA GTGTTATCAC TCATGGTTAT GGCAGCACTG
AGTCTTCATT CAACCGGCCT CACAATAGTG AGTACCAATA CCGTCGTGAC

7701 CATAATTCTC TTACTGTCAT GCCATCCGTA AGATGCTTTT CTGTGACTGG
GTATTAAGAG AATGACAGTA CGGTAGGCAT TCTACGAAAA GACACTGACC

7751 TGAGTACTCA ACCAAGTCAT TCTGAGAATA GTGTATGCAG CGACCGAGTT
ACTCATGAGT TGGTTCAGTA AGACTCTTAT CACATACGCC GCTGGCTCAA

7801 GCTCTGCCG GGCAGTCAATA CGGGATAATA CCGCGCCACA TAGCAGAACT
CGAGAACGGG CCGCAGTTAT GCCCTATTAT GGCGCGGTGT ATCGTCTTGA

7851 TTAAAAGTGC TCATCATTGG AAAACGTTCT TCGGGGCGAA AACTCTCAAG
AATTTCACG AGTAGTAACC TTTTGAAGA AGCCCCGCTT TTGAGAGTTC

7901 GATCTTACCG CTGTTGAGAT CCAGTTCGAT GTAACCCACT CGTGCACCCA
CTAGAATGGC GACAACCTCTA GGTCAAGCTA CATTGGGTGA GCACGTGGGT

7951 ACTGATCTTC AGCATTTTT ACTTTCACCA GCGTTTCTGG GTGAGCAAAA
TGACTAGAAG TCGTAGAAAA TGAAAGTGGT CGCAAAGACC CACTCGTTT

8001 ACAGGAAGGC AAAATGCCGC AAAAAAGGGA ATAAGGGCGA CACGGAAATG
TGTCCCTCCG TTTTACGGCG TTTTTCCCT TATTCCCGCT GTGCCTTTAC

8051 TTGAATACTC ATACTCTTCC TTTTCAATA TTATTGAAGC ATTTATCAGG
AACTTATGAG TATGAGAAGG AAAAAGTTAT AATAACTTCG TAAATAGTCC

8101 GTTATTGTCT CATGAGCGGA TACATATTTG AATGTATTTA GAAAAATAAA
CAATAACAGA GTACTCGCCT ATGTATAAAC TTACATAAAAT CTTTTATTT

8151 CAAATAGGGG TTCCGCGCAC ATTTC
GTTTATCCCC AAGGCGCGTG TAAAG

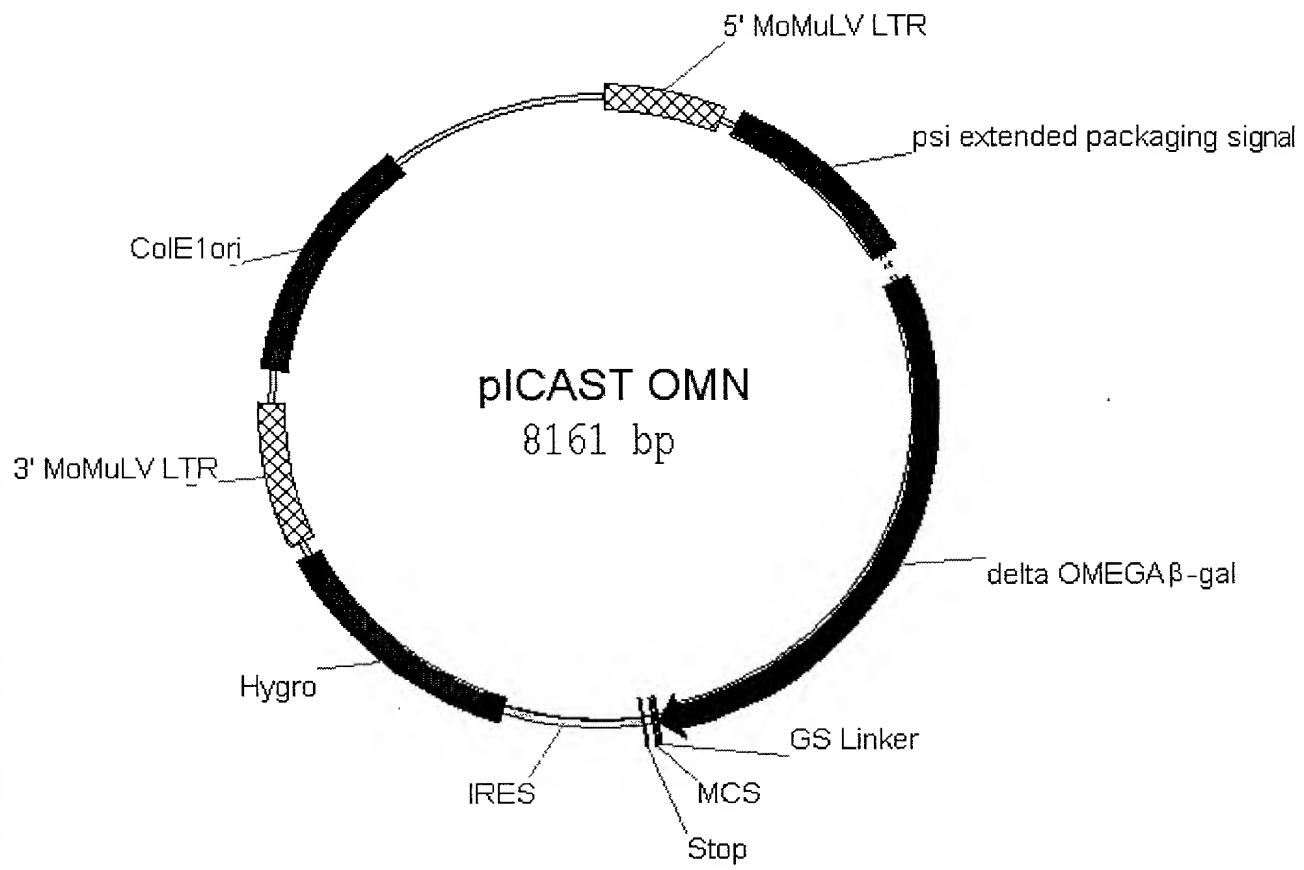


Figure 13A

1 CTGCAGCCTG AATATGGGCC AAACAGGATA TCTGTGGTAA GCAGTTCCTG
GACGTCGGAC TTATACCCGG TTTGTCTAT AGACACCATT CGTCAAGGAC

51 CCCCCGGCTCA GGGCCAAGAAA CAGATGGAAC AGCTGAATAT GGGCCAAACA
GGGGCCGAGT CCCGGTTCTT GTCTACCTTG TCGACTTATA CCCGGTTGT

101 GGATATCTGT GTTAAGCAGT TCCTGCCCG GCTCAGGGCC AAGAACAGAT
CCTATAGACA CCATTCTGCA AGGACGGGGC CGAGTCCCAG TTCTTGTCTA

151 GGTCCCCAGA TCGGGTCCAG CCCTCAGCAG TTTCTAGAGA ACCATCAGAT
CCAGGGGTCT ACGCCAGGTC GGGAGTCGTC AAAGATCTCT TGCTAGTCTA

201 GTTTCCAGGG TGCCCCAAGG ACCTGAAATG ACCCTGTGCC TTATTTGAAC
CAAAGGTCCC ACGGGGTTCC TGGACTTAC TGGGACACGG AATAAACTTG

251 TAACCAATCA GTTCGCTTCT CGCTTCTGTT CGCGCGCTTC TGCTCCCCGA
ATTGGTTAGT CAAGCGAAGA GCGAAGACAA GCGCGCGAAG ACGAGGGCT

301 GCTCAATAAA AGAGCCCACA ACCCCTCACT CGGGGCGCCA GTCCTCCGAT
CGAGTTATTT TCTCGGGTGT TGGGAGTGA GCCCGCGGT CAGGAGGCTA

351 TGACTGAGTC GCCCGGGTAC CCGTGTATCC AATAAACCT CTTGCAGTTG
ACTGACTCAG CGGGCCCATG GGCACATAGG TTATTTGGGA GAACGTCAAC

401 CATCCGACTT GTGGTCTCGC TGTTCCCTGG GAGGGTCTCC TCTGAGTGAT
GTAGGCTGAA CACCAAGAGCG ACAAGGAACC CTCCCAGAGG AGACTCACTA

451 TGACTACCCG TCAGCGGGGG TCTTTCATTT GGGGGCTCGT CCGGGATCGG
ACTGATGGGC AGTCGCCCCC AGAAAGTAAA CCCCCGAGCA GGCCCTAGCC

501 GAGACCCCTG CCCAGGGACC ACCGACCCAC CACCGGGAGG CAAGCTGGCC
CTCTGGGAC GGGTCCCTGG TGGCTGGGTG GTGGCCCTCC GTTCGACCGG

551 ACCAACTTAT CTGTGTCTGT CCGATTGTCT AGTGTCTATG ACTGATTTA
TCGTTGAATA GACACAGACA GGCTAACAGA TCACAGATAC TGACTAAAAT

601 TGCGCCTGCG TCGGTACTAG TTAGCTAACT AGCTCTGTAT CTGGCGGACC
ACGCGGACGC AGCCATGATC AATCGATTGA TCGAGACATA GACCGCCTGG

651 CGTGGTGGAA CTGACGAGTT CTGAACACCC GGCGCGCAACC CTGGGAGACG
GCACCACCTT GACTGCTCAA GACTTGTGGG CCAGCGTTGG GACCCTCTGC

701 TCCCAGGGAC TTTGGGGGCC GTTTTGTGG CCCGACCTGA GGAAGGGAGT
AGGGTCCCTG AAACCCCCGG CAAAAACACC GGGCTGGACT CCTTCCCTCA

751 CGATGTGGAA TCCGACCCCG TCAGGATATG TGGTTCTGGT AGGAGACGAG
GCTACACCTT AGGCTGGGC AGTCCTATAC ACCAAGACCA TCCTCTGCTC

801 AACCTAAAAC AGTTCCCGCC TCCGCTCTGAA TTTTGCTTT CGGTTGGAA
TTGGATTGGTCAAGGGCGG AGGCAGACTT AAAAACGAAA GCCAAACCTT

851 CCGAAGCCGC GCGCTTGTGTC TGCTGCAGCA TCGTTCTGTG TTGTCTCTGT
GGCTTCGGCG CGCAGAACAG ACGACGTCGT AGCAAGACAC AACAGAGACA

901 CTGACTGTGT TTCTGTATTT GTCTGAAAAT TAGGGCCAGA CTGTTACAC
GACTGACACA AAGACATAAA CAGACTTTA ATCCCGGTCT GACAATGGTG

FIGURE 13B

951 TCCCTTAAGT TTGACCTTAG GTAACTGGAA AGATGTCGAG CGGCTCGCTC
AGGAATTCA AACTGGAATC CATTGACCTT TCTACAGCTC GCCGAGCGAG

1001 ACAACCAGTC GGTAGATGTC AAGAAGAGAC GTTGGGTTAC CTTCTGCTCT
TGTTGGTCAG CCATCTACAG TTCTTCTCTG CAACCCAATG GAAGACGGAGA

1051 GCAGAATGGC CAACCTTTAA CGTCGGATGG CGCGGAGACG GCACCTTTAA
CGTCTTACCG GTTGGAAATT GCAGCCTACC GGCGCTCTGC CGTGGAAATT

1101 CCGAGACCTC ATCACCCAGG TTAAGATCAA GGTCTTTCA CCTGGCCC
GGCTCTGGAG TAGTGGGTCC AATTCTAGTT CCAGAAAAGT GGACCGGGCG

1151 ATGGACACCC AGACCAGGTC CCCTACATCG TGACCTGGGA AGCCTGGCT
TACCTGTGGG TCTGGTCCAG GGGATGTAGC ACTGGACCCCT TCGGAACCGA

1201 TTTGACCCCCC CTCCCTGGGT CAAGCCCTTT GTACACCCTA AGCCTCCGCC
AAACTGGGGG GAGGGACCCA GTTCGGGAA CATGTGGGAT TCGGAGGCCG

1251 TCCTCTTCCT CCATCCGCC CGTCTCTCCC CCTGAACCT CCTCGTCGA
AGGAGAAGGA GGTAGGCGGG GCAGAGAGGG GGAACCTTGA GGAGCAAGCT

1301 CCCCCCCTCG ATCCCTCCCT TATCCAGCCC TCACTCCTTC TCTAGGCGCC
GGGGCGGAGC TAGGAGGGAA ATAGGTGGG AGTGAGGAAG AGATCCCGG

1351 GGGCGCTCTA GCCCATTAAT ACGACTCACT ATAGGGCGAT TCGAACACCA
CCGGCGAGAT CGGGTAATTA TGCTGAGTGA TATCCCGCTA AGCTTGTGGT

1401 TGCACCATCA TCATCATCAC GTCGACGAAC AGAAACTCAT TTCCGAAGAA
ACGTGGTAGT AGTAGTAGTG CAGCTGCTTG TCTTGAGTA AAGGCTTCTT

1451 GACCTACTCG AGATGGGCGT GATTACGGAT TCACTGGCCG TCGTTTACA
CTGGATGAGC TCTACCCGCA CTAATGCCTA AGTGACCGGC AGCAAAATGT

1501 ACGTCGTGAC TGGGAAACCC CTGGCGTTAC CCAACTTAAT CGCCTTGCAG
TGCAGCACTG ACCCTTTGG GACCGCAATG GGTTGAATTA GCGGAACGTC

1551 CACATCCCCC TTCGCCAGC TGGCGTAATA GCGAAGAGGC CGCACCAGAT
GTGTAGGGGG AAAGCGGTG ACCGCATTAT CGCTTCTCCG GCGGTGGCTA

1601 CGCCCTTCCC AACAGTTACG CAGCCTGAAT GCGAATGGC GCTTGCCTG
CGGGGAAGGG TTGTCAATGC GTCGGACTTA CCGCTTACCG CGAACCGAC

1651 GTTTCGGCA CCAGAAGCGG TGCCGGAAAG CTGGCTGGAG TCGGATCTTC
CAAAGGCCGT GGTCTCGCC ACGGCCTTC GACCGACCTC ACGCTAGAAG

1701 CTGAGGCCGA TACTGTCGTC GTCCCCCTCAA ACTGGCAGAT GCACGGTTAC
GACTCCGGCT ATGACAGCAG CAGGGAGTT TGACCGTCTA CGTGCCAATG

1751 GATGCGCCCA TCTACACCAA CGTGACCTAT CCCATTACGG TCAATCCGCC
CTACGGGGT AGATGTGGTT GCACTGGATA GGGTAATGCC AGTTAGGCAG

1801 GTTTGTTCCC ACGGAGAACG CGACGGGTTG TTACTCGCTC ACATTAATG
CAAACAAGGG TGCCTCTTAG GCTGCCAAC AATGAGCGAG TGTAAATTAC

1851 TTGATGAAAG CTGGCTACAG GAAGGCCAGA CGCGAATTAT TTTTGATGGC
AACTACTTTC GACCGATGTC CTTCCGGTCT GCGCTTAATA AAAACTACCG

1901 GTTAACTCGG CGTTTCATCT GTGGTGCAAC GGGCGCTGGG TCGGTTACGG
CAATTGAGCC GCAAAGTAGA CACCACGTTG CCCGCGACCC AGCCAATGCC

1951 CCAGGACAGT CGTTGCCGT CTGAATTGGA CCTGAGCGCA TTTTACGCG
GGTCCTGTCA GCAAACGGCA GACTTAAACT GGACTCGCGT AAAAATGCGC

2001 CCGGAGAAAA CCGCCTCGCG GTGATGGTGC TGCGCTGGAG TGACGGCAGT
GGCCTCTTT GGCGGAGCGC CACTACCACG ACGCGACCTC ACTGCCGTCA

2051 TATCTGGAAG ATCAGGATAT GTGGCGGATG AGCGGCATT TCCGTGACGT
ATAGACCTTC TAGTCCTATA CACCGCCTAC TCGCCGTAAA AGGCACGTCA

2101 CTCGTTGCTG CATAAACCGA CTACACAAAT CAGCGATTTC CATGTTGCCA
GAGCAACGAC GTATTGGCT GATGTGTTA GTCGCTAAAG GTACAACGGT

2151 CTCGTTAA TGATGATTTC AGCCCGCCTG TACTGGAGGC TGAAGTTCAG
GAGCGAAATT ACTACTAAAG TCGGCGCGAC ATGACCTCCG ACTTCAAGTC

2201 ATGTGCGGCG AGTTGCGTGA CTACCTACGG GTAACAGTTT CTTTATGGCA
TACACGCCGC TCAACGCACT GATGGATGCC CATTGTCAAA GAAATACCGT

2251 GGGTGAACG CAGGTGCCA GCGGCACCGC GCCTTCGGC GGTGAAATTA
CCCACTTGC GTCCAGCGGT CGCCGTGGCG CGGAAAGCGC CCACTTAAT

2301 TCGATGAGCG TGGTGGTTAT GCGATCGCG TCACACTACG TCTGAACGTC
AGCTACTCGC ACCACCAATA CGGCTAGCGC AGTGTGATGC AGACTTGCAG

2351 GAAAACCGA AACTGTGGAG CGCCGAAATC CGAAATCTCT ATCGTGCCTG
CTTTGGGCT TTGACACCTC GCGGCTTAG GGCTTAGAGA TAGCACGCCA

2401 GGTTGAAC TG CACACGCCG ACGGCACGCT GAATTGAAGCA GAAGCCTGCG
CCAACTTGAC GTGTGGCGC TGCGTGCCTA CTAACCTCGT CTTCGGACGC

2451 ATGTCGGTTT CCGCGAGGTG CGGATTGAAA ATGGTCTGCT GCTGCTGAAC
TACAGCCAAA GGCCTCCAC GCCTAACTTT TACCAAGACGA CGACGACTTG

2501 GGCAAGCCGT TGCTGATTG AGGCCTTAAC CGTCACGAGC ATCATCCTCT
CCGTCGGCA ACGACTAAGC TCCGCAATTG GCAGTGCTCG TAGTAGGAGA

2551 GCATGGTCAG GTCATGGATG AGCAGACGAT GGTGCAGGAT ATCCTGCTGA
CGTACCGATC CAGTACCTAC TCGTCTGCTA CCACGTCTA TAGGACGACT

2601 TGAAGCAGAA CAACTTTAAC GCCGTGCGCT GTTCGCTTAA TCCGAACCAT
ACTTCGTCTT GTTGAATTG CGGCACGCGA CAAGCGTAAT AGGCTTGGTA

2651 CCGCTGTGGT ACACGCTGTG CGACCGCTAC GGCCTGTATG TGGTGGATGA
GGCGACACCA TGTGCGACAC GCTGGCGATG CGGGACATAC ACCACCTACT

2701 AGCCAATATT GAAACCCACG GCATGGTGCC AATGAATCGT CTGACCGATG
TCGGTTATAA CTTTGGGTGC CGTACACCGG TTACTTAGCA GACTGGCTAC

2751 ATCCCGCGCTG GCTACCGGGCG ATGAGCGAAC GCGTAACGCG AATGGTGCAG
TAGGCGCGAC CGATGGCCGC TACTCGCTTG CGCATTGCGC TTACCAACGTC

2801 CGCGATCGTA ATCACCCGAG TGTGATCATC TGGTCGCTGG GGAATGAATC
GCGCTAGCAT TAGTGGGCTC ACACTAGTAG ACCAGCGACC CCTTACTTAG

2851 AGGCCACGGC GCTAATCACG ACGCGCTGTA TCGCTGGATC AAATCTGTG
TCCGGTGCCG CGATTAGTGC TGCGCGACAT AGCGACCTAG TTTAGACAGC

2901 ATCCTTCCCG CCCGGTGCAG TATGAAGGCG GCGGAGCCGA CACCACGGCC
TAGGAAGGGC GGGCCACGTC ATACTTCCGC CGCCTCGGCT GTGGTGCCGG

2951 ACCGATATTA TTTGCCCGAT GTACCGCGC GTGGATGAAG ACCAGCCCTT
TGGCTATAAT AACCGGGCTA CATGCCGCG CACCTACTTC TGTCGGAA

3001 CCCGGCTGTG CCGAAATGGT CCATCAAAAAA ATGGCTTTCG CTACCTGGAG
GGGCCGACAC GGCTTTACCA GGTAGTTTT TACCGAAAGC GATGGACCTC

3051 AGACCGCGCCC GCTGATCCTT TGCGAATACG CCCACCGCAT GGGTAACAGT
TCTGCCGCGG CGACTAGGAA ACGCTTATGC GGGTGCCTA CCCATTGTCA

3101 CTTGGCGGTT TCGCTAAATA CTGGCAGGCG TTTCGTCAGT ATCCCCGTTT
GAACCGCAA AGCGATTAT GACCGTCCGC AAAGCAGTCA TAGGGGCAA

3151 ACAGGGCGGC TTCGCTCTGGG ACTGGGTGGA TCAGTCGCTG ATTAAATATG
TGTCCCACCG AAGCAGACCC TGACCCACCT AGTCAGCGAC TAATTATAC

3201 ATGAAAACGG CAACCCGTGG TCGGCTTACG GCGGTGATTG TGGCGATACG
TACTTTGCC GTTGGGCACC AGCCGAATGC CGCCACTAAA ACCGCTATGC

3251 CCGAACGATC GCCAGTTCTG TATGAACGGT CTGGTCTTTG CCGACCGCAC
GGCTTGCTAG CGGTCAAGAC ATACTGCCA GACCAGAAAC GGCTGGCGTG

3301 GCCGCATCCA GCGCTGACGG AAGCAAAACA CCAGCAGCAC TTTTCCAGT
CGGCGTAGGT CGCGACTGCC TTCGTTTGT GGTCGTCGTC AAAAAGGTCA

3351 TCCGTTTATC CGGGCAAACCC ATCGAAGTGA CCAGCGAATA CCTGTTCCGT
AGGCAAATAG GCCCGTTGG TAGCTTCACT GGTCGCTTAT GGACAAGGCA

3401 CATAGCGATA ACGAGCTCCT GCACCTGGATG CTGGCGCTGG ATGGTAAGCC
GTATCGCTAT TGCTCGAGGA CGTGACCTAC CACCGCGACC TACCATTCGG

3451 GCTGGCAAGC GGTGAAGTGC CTCTGGATGT CGCTCCACAA GGTAAACAGT
CGACCGTTCG CCACCTCACG GAGACCTACA GCGAGGGTGT CCATTGTCA

3501 TGATTGAACG GCCTGAACTA CCGCAGCCGG AGAGCGCCGG GCAACTCTGG
ACTAACTTGA CGGACTTGAT GGCCTCGGCC TCTCGCGGCC CGTTGAGACC

3551 CTCACAGTAC GCGTAGTGCA ACCGAACGCG ACCGCATGGT CAGAAGCCGG
GAGTGTCTAG CGCATCACGT TGGCTTGCGC TGGCGTACCA GTCTTCGGCC

3601 GCACATCAGC GCCTGGCAGC AGTGGCGTCT GGCAGGAAAC CTCAGTGTGA
CGTGTAGTCG CGGACCGTCG TCACCCGAGA CGCCCTTTTG GAGTCACACT

3651 CGCTCCCCGC CGCGTCCAC GCCATCCCAGC ATCTGACCCAC CAGCGAAATG
GCGAGGGCG GCGCAGGGTG CGGTAGGGCG TAGACTGGTG GTGCCTTAC

3701 GATTTTGCA TCGAGCTGGG TAATAAGCGT TGGCAATTAA ACCGCCAGTC
CTAAAAACGT AGCTCGACCC ATTATTCGCA ACCGTTAAAT TGGCGGTCA

3751 AGGCTTCTT TCACAGATGT GGATTGGCGA TAAAAAACAA CTGCTGACGC
TCCGAAAGAA AGTGTCTACA CCTAACCGCT ATTGTTTGTT GACGACTGCG

3801 CGCTGCGCGA TCAGTTCACCGTGTGATA GATCTGGAGG TGGTGGCAGC
GCGACGCGCT AGTCAAGTGG GCACAGCTAT CTAGACCTCC ACCACCGTCG

3851 AGGCCTTGGC GCGCCGGATC CTTAATTAAC AATTGACCGG TAATAATAGG
TCCGGAACCG CGCGGCCTAG GAATTAATTG TTAACTGGCC ATTATTATCC

3901 TAGATAACTG ACTGATTAGA TGCAATTGCA CTAGATCCCT CGACCAATTCA
ATCTATTACAC TGACTAATCT ACGTAAAGCT GATCTAGGGGA GCTGGTTAAG

3951 CGGTTATTTT CCACCATATT GCCGTCCTTT GGCAATGTGA GGGCCCCGAA
GCCAATAAAA GGTGGTATAA CGGCAGAAAA CGCTTACACT CCCGGGCCTT

4001 ACCTGGCCCT GTCTTCTTGA CGAGCATTCC TAGGGGTCTT TCCCCTCTCG
TGGACCGGGA CAGAAGAACT GCTCGTAAGG ATCCCCAGAA AGGGGAGAGC

4051 CCAAAGGAAT GCAAGGTCTG TTGAATGTCTG TGAAGGAAGC AGTTCCCTTG
GGTTTCCCTTA CGTCCAGAC AACTTACAGC ACTTCCTCTG TCAAGGAGAC

4101 GAAGCTTCTT GAAGACAAAC AACGTCGTGA GCGACCCCTT GCAGGGCAGCG
CTTCGAAGAA CTTCTGTTTGT TGCAAGACAT CGCTGGGAAA CGTCCGTGCG

4151 GAACCCCCCA CCTGGCGACA GGTGCCTCTG CGGCCAAAAG CCACGTGTAT
CTTGGGGGGT GGACCGCTGT CCACGGAGAC GCCGGTTTC GGTGCACATA

4201 AAGATACACC TGCAAAGGCG GCACAACCCC AGTGCCACGT TGTGAGTTGG
TTCTATGTGG ACGTTCCGC CGTGTGGGG TCACGGTGCA ACACTCAACC

4251 ATAGTTGTGG AAAGAGTCAA ATGGCTCTCC TCAAGCGTAT TCAACAAGGG
TATCAACACC TTTCTCAGTT TACCGAGAGG AGTCGCATA AGTTGTCCC

4301 GCTGAAGGAT GCCCAGAAGG TACCCCATTG TATGGGATCT GATCTGGGC
CGACTTCCTA CGGGTCTTCC ATGGGGTAAC ATACCCCTAGA CTAGACCCCG

4351 CTCGGTGCAC ATGCTTTACA TGTGTTAGT CGAGGTTAAA AAACGTCTAG
GAGCCACGTG TACGAAATGT ACACAAATCA GCTCCAATT TTTGCAGATC

4401 GCCCCCCGAA CCACGGGGAC GTGGTTTCC TTTGAAAAAC ACGATGATAA
CGGGGGGCTT GGTGCCCTG CACCAAAAGG AAACTTTTG TGCTACTATT

4451 TACCATGAAA AAGCCTGAAC TCACCGCGAC GTCTGTCGAG AAGTTCTGA
ATGGTACTTT TTCGGACTTG AGTGGCGCTG CAGACAGCTC TTCAAAGACT

4501 TCGAAAAGTT CGACAGCGTC TCCGACCTGA TGCAGCTCTC GGAGGGCGAA
AGCTTTCAA GCTGTCGAG AGGCTGGACT ACGTCGAGAG CCTCCCGCTT

4551 GAATCTCGTG CTTTCAGCTT CGATGTAGGA GGGCGTGGAT ATGTCCTGCG
CTTAGAGCAC GAAAGTCGAA GCTACATCCT CCCGCACCTA TACAGGACGC

4601 GGTAAATAGC TCGCGCGATG GTTCTACAA AGATCGTTAT GTTTATCGGC
CCATTATCG ACACGGCTAC CAAAGATGTT TCTAGCAATA CAAATAGCCG

4651 ACTTTGCATC GGCGCGCTC CCGATTCCGG AAGTGCTTGA CATTGGGGAA
TGAAACGTAG CGGGCGCGAG GGCTAAGGCC TTCACGAACCTT GTAACCCCTT

4701 TTTAGCGAGA GCCTGACCTA TTGCATCTCC CGCCGTGCAC AGGGTGTAC
AAATCGCTCT CGGACTGGAT AACGTAGAGG GCGGCACGTG TCCCACAGTG

4751 GTTGAAGAC CTGCCTGAAA CGAACTGCC CGCTGTTCTG CAGCCGGTCG
CAACGTTCTG GACGGACTTT GGCTTGACGG GCGACAAGAC GTCGGCCAGC

4801 CGGAGGCCAT GGATGCGATC GCTGCCGCCG ATCTTAGCCA GACGAGCGGG
GCCTCCGGTA CCTACGCTAG CGACGCCGGC TAGAATCGGT CTGCTGCC

4851 TTCCGGCCAT TCGGACCGCA AGGAATCGGT CAATACACTA CATGGCGTGA
AAGCCGGGTAGCCTAGCGT TCCTTAGCCA GTTATGTGAT GTACCGCACT

4901 TTTCATATGC GCGATTGCTG ATCCCCATGT GTATCACTGG CAAACTGTGA
AAAGTATAACG CGCTAACGAC TAGGGGTACA CATAGTGACC GTTTGACACT

4951 TGGACGACAC CGTCAGTGC TCCGTCGCG AGGCTCTCGA TGAGCTGATG
ACCTGCTGTG GCAGTCACGC AGGCAGCGCG TCCGAGAGCT ACTCGACTAC

5001 CTTTGGGCCG AGGACTGCC CGAAGTCCGG CACCTCGTGC ACGCGGATTT
GAAACCCGGC TCCTGACGGG GCTTCAGGCC GTGGAGCACG TGCGCCTAAA

5051 CGGCTCCAAC AATGTCTGA CGGACAATGG CGCGATAACA GCGGTCAATTG
GCCGAGGTTG TTACAGGACT GCCTGTTACC GGCGTATTGT CGCCAGTAAC

5101 ACTGGAGCGA GGCGATGTTG GGGGATTCCC AATACGAGGT CGCCAACATC
TGACCTCGCT CCGCTACAAG CCCCTAAGGG TTATGCTCCA GCGGTTGTAG

5151 TTCTTCTGGA GGCGTGGTT GGCTTGTATG GAGCAGCAGA CGCGCTACTT
AAGAAGACCT CCGGCACCAA CGAACATAC CTCGTCGTCT GCGCGATGAA

5201 CGAGCGGAGG CATCCGGAGC TTGCAGGATC GCCGCGGCTC CGGGCGTATA
GCTCGCCTCC GTAGGCCTCG AACGTCCTAG CGGCGCCGAG GCCCAGCATAT

5251 TGCTCCGCAT TGGTCTTGAC CAACTCTATC AGAGCTTGGT TGACGGCAAT
ACGAGGCGTA ACCAGAACTG GTTGAGATAG TCTCGAACCA ACTGCCGTTA

5301 TTCGATGATG CAGCTGGGC GCAGGGTCGA TGCGACGCAA TCGTCCGATC
AAGCTACTAC GTCGAACCCG CGTCCAGCT ACGCTGCGTT AGCAGGCTAG

5351 CGGAGCCGGG ACTGTCGGGC GTACACAAAT CGCCCGCAGA AGCGCGGCCG
GCCTCGGCCG TGACAGCCCG CATGTGTTA GCGGGCGTCT TCGCGCCGGC

5401 TCTGGACCGA TGGCTGTGTA GAAGTACTCG CCGATAGTGG AAACCGACGC
AGACCTGGCT ACCGACACAT CTTCATGAGC GGCTATCACC TTTGGCTGCG

5451 CCCAGCACTC GTCCGAGGGC AAAGGAATAG AGTAGATGCC GACCAGGATC
GGGTCGTGAG CAGGCTCCCG TTTCCTTATC TCATCTACGG CTGGCCCTAG

5501 TATCGATAAA ATAAAAGATT TTATTTAGTC TCCAGAAAAA GGGGGGAATG
ATAGCTATTT TATTTCTAA AATAAATCAG AGGTCTTTTT CCCCCCTTAC

5551 AAAGACCCCA CCTGTAGGTT TGGCAAGCTA GCTTAAGTAA CGCCATTG
TTCTGGGGT GGACATCCAA ACCGTTCGAT CGAATTCAATT GCGGTAAAAC

5601 CAAGGCATGG AAAAATACAT AACTGAGAAT AGAGAAGTTC AGATCAAGGT
GTTCCGTACC TTTTATGTA TTGACTCTTA TCTCTTCAAG TCTAGTTCCA

5651 CAGGAACAGA TGGAACAGCT GAATATGGGC CAAACAGGAT ATCTGTGGTA
GTCCTTGTCT ACCTTGTGCA CTTATACCCG GTTTGTCCCTA TAGACACCAT

5701 AGCAGTTCCCT GCCCCGGCTC AGGGCCAAGA ACAGATGGAA CAGCTGAATA
TCGTCAGGA CGGGGCCGAG TCCCGBTCT TGTCTACCTT GTCGACTTAT

5751 TGGGCCAAAC AGGATATCTG TGGTAAGCAG TTCTGCCCG GGCTCAGGGC
ACCCGGTTG TCCTATAGAC ACCATCGTC AAGGACGGGG CCGAGTCCCG

5801 CAAGAACAGA TGGTCCCCAG ATGCCGTCCA GCCCTCAGCA GTTTCTAGAG
GTTCTGTCT ACCAGGGGTAC TACGCCAGGT CGGGAGTCGT CAAAGATCTC

5851 AACCATCAGA TGTTTCCAGG GTGCCCAAG GACCTGAAAT GACCCTGTGC
TTGGTAGTCT ACAAAAGGTCC CACGGGGTTC CTGGACTTTA CTGGGACACG

5901 CTTATTTGAA CTAACCAATC AGTTCGCTTC TCGCTTCTGT TCGCGCGCTT
GAATAAACTT GATTGGTTAG TCAAGCGAAG AGCGAAGACA AGCGCGCGAA

5951 CTGCTCCCCG AGCTCAATAA AAGAGCCCAC AACCCCTCAC TCAGGGCGCC
GACGAGGGC TCGAGTTATT TTCTCGGGTG TTGGGGAGTG AGCCCGCGGG

6001 AGTCCTCCGA TTGACTGAGT CGCCCGGGTA CCCGTGTATC CAATAAACCC
TCAGGAGGCT AACTGACTCA GCGGGCCCCAT GGGCACATAG GTTATTGGG

6051 TCTTGCAGTT GCATCCGACT TGTGGTCTCG CTGTTCTTG GGAGGGCTC
AGAACGTCAA CGTAGGCTGA ACACCAGAGC GACAAGGAAC CCTCCAGAG

6101 CTCTGAGTGA TTGACTACCC GTCAGCGGGG GTCTTTCATT CATGCAGCAT
GAGACTCACT AACTGATGGG CAGTCGCCCC CAGAAAAGTAA GTACGTCGT

6151 GTATCAAAT TAATTTGGTT TTTTTCTTA AGTATTACAA TTAAATGGCC
CATAGTTTA ATTAAACCAA AAAAAGAAT TCATAATGT AATTACCGG

6201 ATAGTTGCAT TAATGAATCG GCCAACGCGC GGGGAGAGGC GGTTTGCCTA
TATCAACGTA ATTACTTAGC CGGTTGCGCG CCCCTCTCCG CCAAACGCAT

6251 TTGGCGCTCT TCCGCTTCT CGCTCACTGA CTCGCTGCGC TCGGTGTT
AACCGCGAGA AGGCGAAGGA GCGAGTGACT GAGCGACGCG AGCCAGCAAG

6301 GGCTGCGGCG AGCGGTATCA GCTCACTCAA AGGCGGTAAT ACGGTTATCC
CCGACGCCGC TCGCCATAGT CGAGTGAGTT TCCGCCATTA TGCCAATAGG

6351 ACAGAACATCG GGGATAACGC AGGAAAGAAC ATGTGAGCAA AAGGCCAGCA
TGTCTTAGTC CCCTATTGCG TCCTTCTTG TACACTCGTT TTCCGGTCGT

6401 AAAGGCCAGG AACCGTAAAA AGGCCCGGTT GCTGGCGTT TTCCATAGGC
TTCCGGTCC TTGGCATTTC TCCGGCGAA CGACCGCAA AAGGTATCCG

6451 TCCGGCCCCC TGACGAGCAT CACAAAAATC GACGCTCAAG TCAGAGGTGG
AGGCGGGGG ACTGCTCGTA GTGTTTTAG CTGCGAGTTC AGTCTCCACC

6501 CGAAACCCGA CAGGACTATA AAGATACCAAG GCGTTCCCC CTGGAAAGCTC
GCTTGGGCT GTCCTGATAAT TTCTATGGTC CGCAAAGGGG GACCTTCGAG

6551 CCTCGTGCAGC TCTCCTGTT CGACCCCTGCC GCTTACCGGA TACCTGTCCG
GGAGCACGCG AGAGGACAAG GCTGGGACGG CGAATGGCCT ATGGACAGGC

6601 CCTTCTCCC TTGGGAAGC GTGGCGCTTT CTCATAGCTC ACGCTGTAGG
GGAAAGAGGG AAGCCCTTCG CACCGCGAAA GAGTATCGAG TGCGACATCC

6651 TATCTCAGTT CGGTGTAGGT CGTCGCTCC AAGCTGGGCT GTGTGCACGA
ATAGAGTCAA GCCACATCCA GCAAGCGAGG TTCGACCCGA CACACGTGCT

6701 ACCCCCCGTT CAGCCCGACC GCTGCCCTT ATCCGGTAAC TATCGTCTTG
TGGGGGGCAA GTCGGGCTGG CGACCGGAA TAGGCCATTG ATAGCAGAAC

6751 AGTCCAACCC GGTAAGACAC GACTTATCGC CACTGGCAGC AGCCACTGGT
TCAGGTTGGG CCATTCTGTG CTGAATAGCG GTGACCGTCG TCGGTGACCA

6801 AACAGGATTA GCAGAGCGAG GTATGTAGGC GGTGCTACAG AGTTCTGAA
TTGTCTTAAT CGTCTCGCTC CATAACATCCG CCACGATGTC TCAAGAACTT

6851 GTGGTGGCCT AACTACGGCT ACACAGAAG AACAGTATTG GGTATCTGCG
CACCAACCGGA TTGATGCCGA TGTGATCTTC TTGTCATAAA CCATAGACGC

6901 CTCTGCTGAA GCCAGTTACC TTGGAAAAAA GAGTTGGTAG CTCTTGATCC
GAGACGACTT CGGTCAATGG AAGCCTTTT CTCAACCACATC GAGAACTAGG

6951 GGCAAAACAAA CCACCGCTGG TAGCGGTGGT TTTTTGTTT GCAAGCAGCA
CCGTTTGTGTT GGTGGCGACC ATCGCCACCA AAAAAACAAA CGTTCGTCGT

7001 GATTACGCGC AGAAAAAAAG GATCTCAAGA AGATCCTTTG ATCTTTCTA
CTAATGCGCG TCTTTTTTC CTAGAGTTCT TCTAGGAAAC TAGAAAAGAT

7051 CGGGGTCTGA CGCTCAGTGG AACGAAAAGT CACGTTAAGG GATTTGGTC
GCCCGAGACT GCGAGTCACC TTGCTTTGA GTGCAATTCC CTAAACCCAG

7101 ATGAGATTAT CAAAAAGGAT CTTCACCTAG ATCCTTTGC GGCGCAAAT
TACTCTAATA GTTTTCCTA GAAGTGGATC TAGGAAAACG CCGCGTTA

7151 CAATCTAAAG TATATATGAG TAAACTTGGT CTGACAGTTA CCAATGCTTA
GTTAGATTTC ATATATACTC ATTTGAACCA GACTGTCAAT GGTTACGAAT

7201 ATCAGTGAGG CACCTATCTC AGCGATCTGT CTATTTCGTT CATCCATAGT
TAGTCACTCC GTGGATAGAG TCGCTAGACA GATAAAGCAA GTAGGTATCA

7251 TGCTGACTC CCCGTCGTGT AGATAACTAC GATACGGGAG GGCTTACCAT
ACGGACTGAG GGGCAGCACA TCTATTGATG CTATGCCCTC CCGAATGGTA

7301 CTGGCCCCAG TGCTGCAATG ATACCGCGAG ACCCACGCTC ACCGGCTCCA
GACCGGGGTC ACACGTTAC TATGGCGCTC TGGGTGCGAG TGGCCGAGGT

7351 GATTATTCAG CAATAAACCA GCCAGCCGGA AGGGCCGAGC GCAGAAGTGG
CTAAATAGTC GTTATTGGT CGGTCGGCCT TCCCAGCTCG CGTCTTCACC

7401 TCCTGCAACT TTATCCGCT CCATCCAGTC TATTAATTGT TGCCGGGAAG
AGGACGTTGA AATAGGCGGA GGTAGGTCAG ATAATTAACA ACGGCCCTTC

7451 CTAGAGTAAG TAGTTGCCA GTTAATAGTT TGCACAGT TGTTGCCATT
GATCTCATTC ATCAAGCGGT CAATTATCAA ACGCGTTGCA ACAACGGTAA

7501 GCTACAGGCA TCGTGGTGTAC ACGCTCGTCG TTTGGTATGG CTTCATTCA
CGATGTCCGT AGCACCAACAG TGCAGACAGC AAACCATACC GAAGTAAGTC

7551 CTCCGGTTCC CAACGATCAA GGCGAGTTAC ATGATCCCCC ATGTTGTGCA
GAGGCCAAGG GTTGCTAGTT CCGCTCAATG TACTAGGGGG TACAACACGT

7601 AAAAAGCGGT TAGCTCCTTC GGTCCCTCCGA TCGTTGTCAG AAGTAAGTTG
TTTTTCGCCA ATCGAGGAAG CCAGGAGGCT AGCAACAGTC TTCATTCAAC

7651 GCCGCAGTGT TATCACTCAT GGTTATGGCA GCACTGCATA ATTCTCTTAC
CGGCCTCACCA ATAGTGAGTA CCAATACCGT CGTGACGTAT TAAGAGAAATG

7701 TGTCATGCCA TCCGTAAGAT GCTTTCTGT GACTGGTGAG TACTCAACCA
ACAGTACGGT AGGCATTCTA CGAAAAGACA CTGACCACTC ATGAGTTGGT

7751 AGTCATTCTG AGAATAGTGT ATGCGGCGAC CGAGTTGCTC TTGCCCCGCG
TCAGTAAGAC TCTTATCACA TACGCCGCTG GCTCAACGAG AACGGGCCGC

7801 TCAATACGGG ATAATACCGC GCCACATAGC AGAACCTTAA AAGTGCTCAT
AGTTATGCCC TATTATGGCG CGGTGTATCG TCTTGAATT TTCACGAGTA

7851 CATTGGAAAA CGTTCTTCGG GGCAGAAACT CTCAGGATC TTACCGCTGT
GTAACCTTTT GCAAGAAGCC CCGCTTTGA GAGTTCTAG AATGGCGACA

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8001 TGCCGCAAAA AAGGGAATAA GGGCGACACG GAAATGTTGA ATACTCATAC
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8051 TCTTCCTTTT TCAATATTAT TGAAGCATTG ATCAGGGTTA TTGTCTCATG
AGAAGGAAAA AGTTATAATA ACTTCGTAAA TAGTCCAAT AACAGAGTAC

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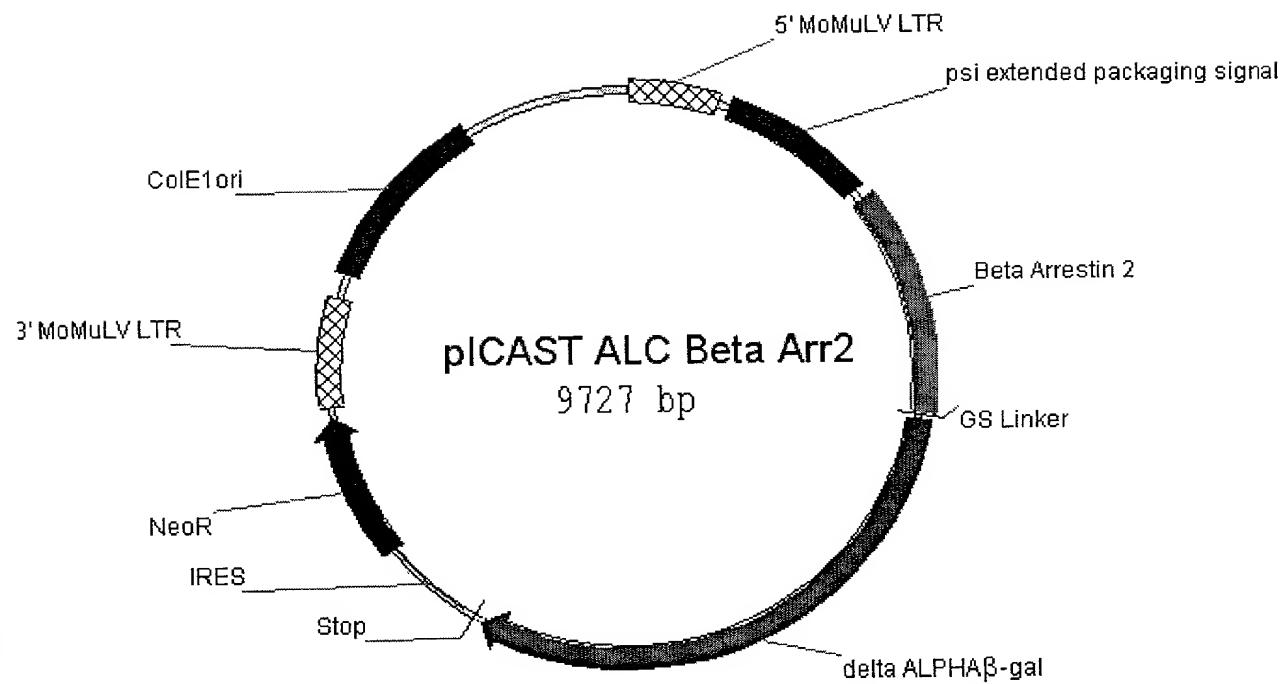


Figure 14

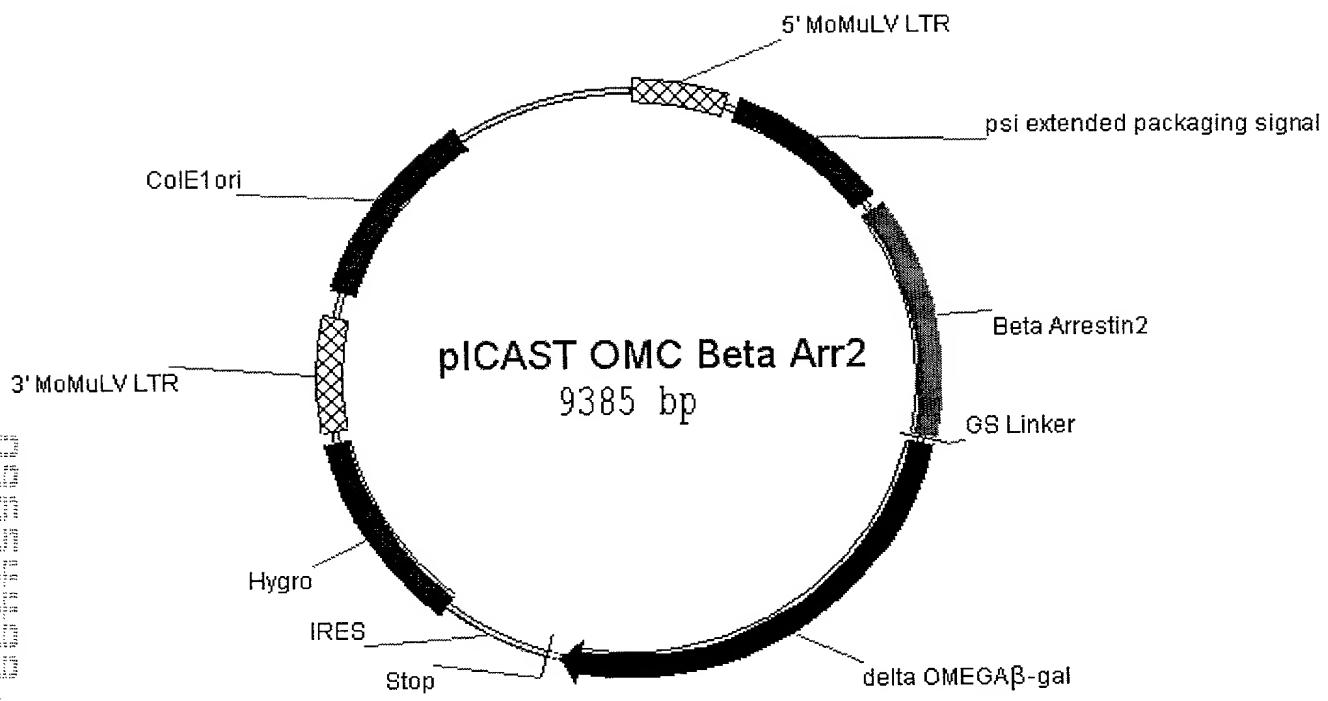


Figure 15

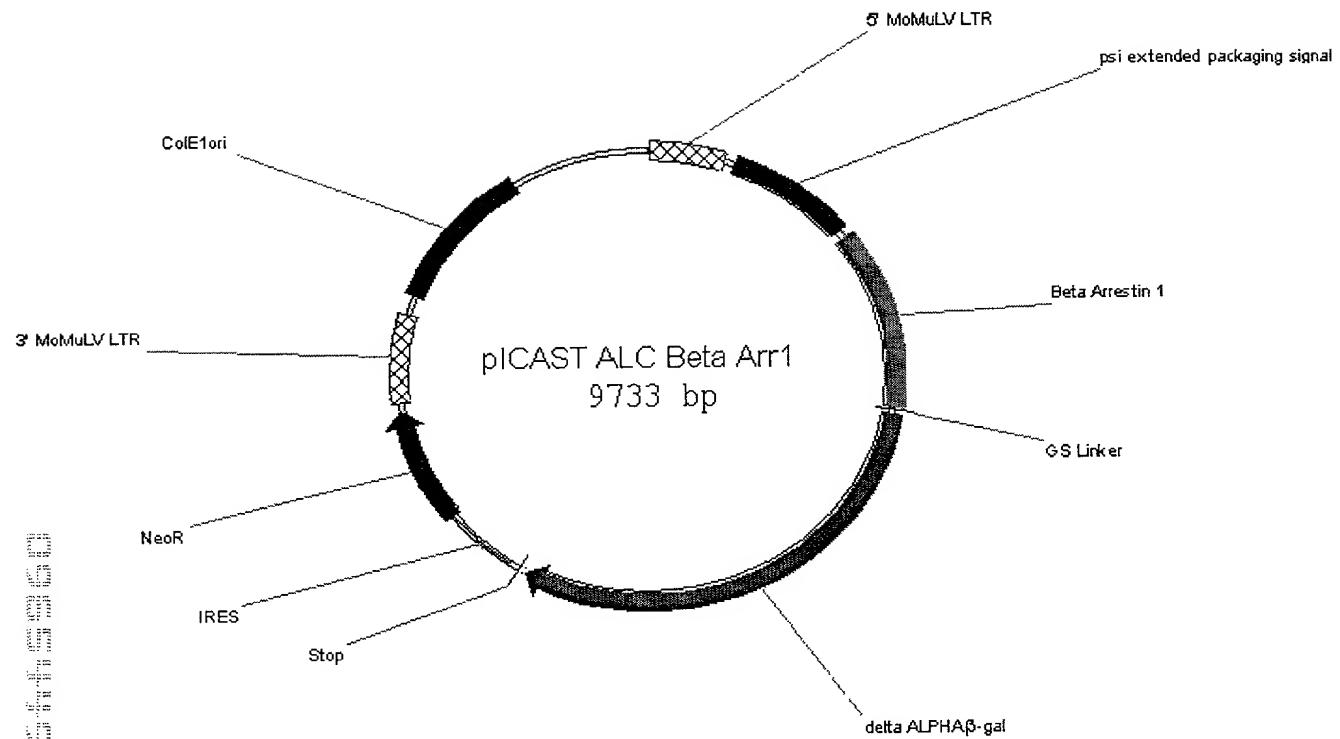


Figure 16

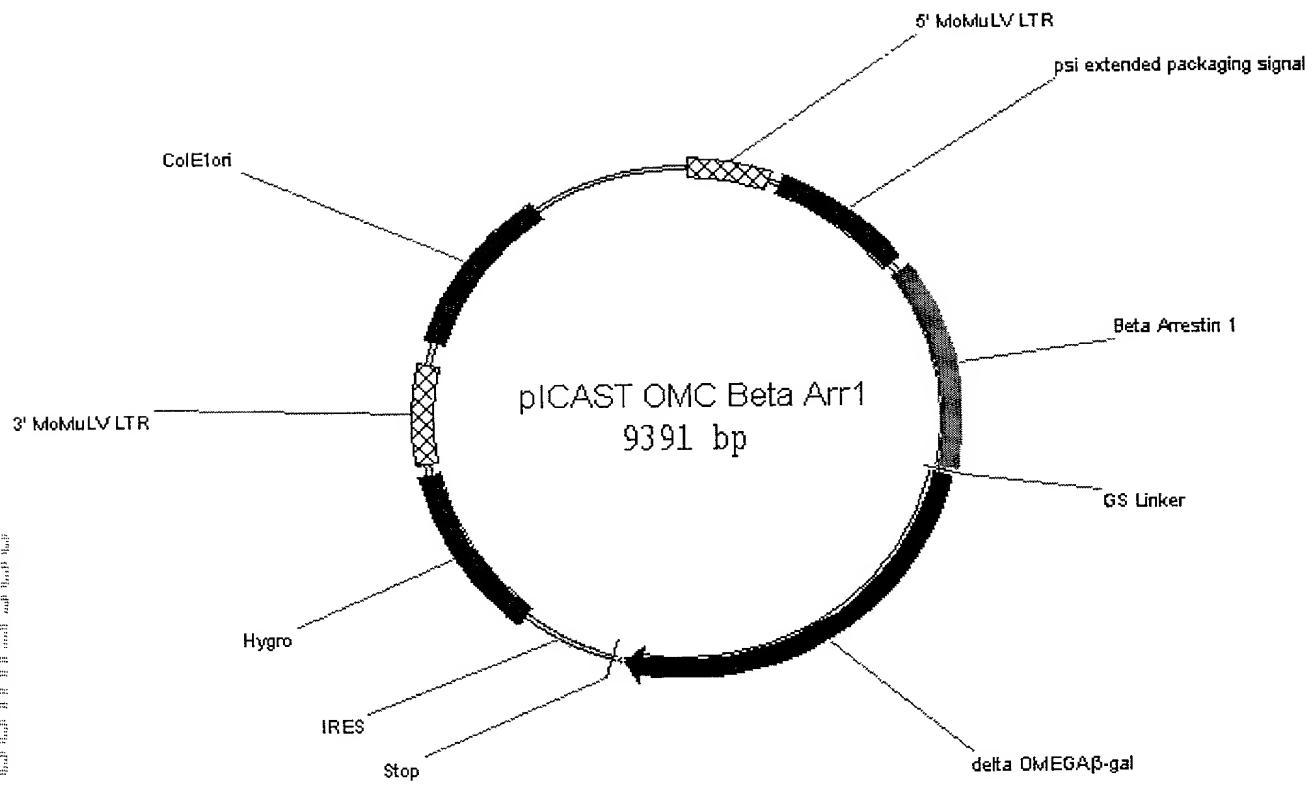


Figure 17

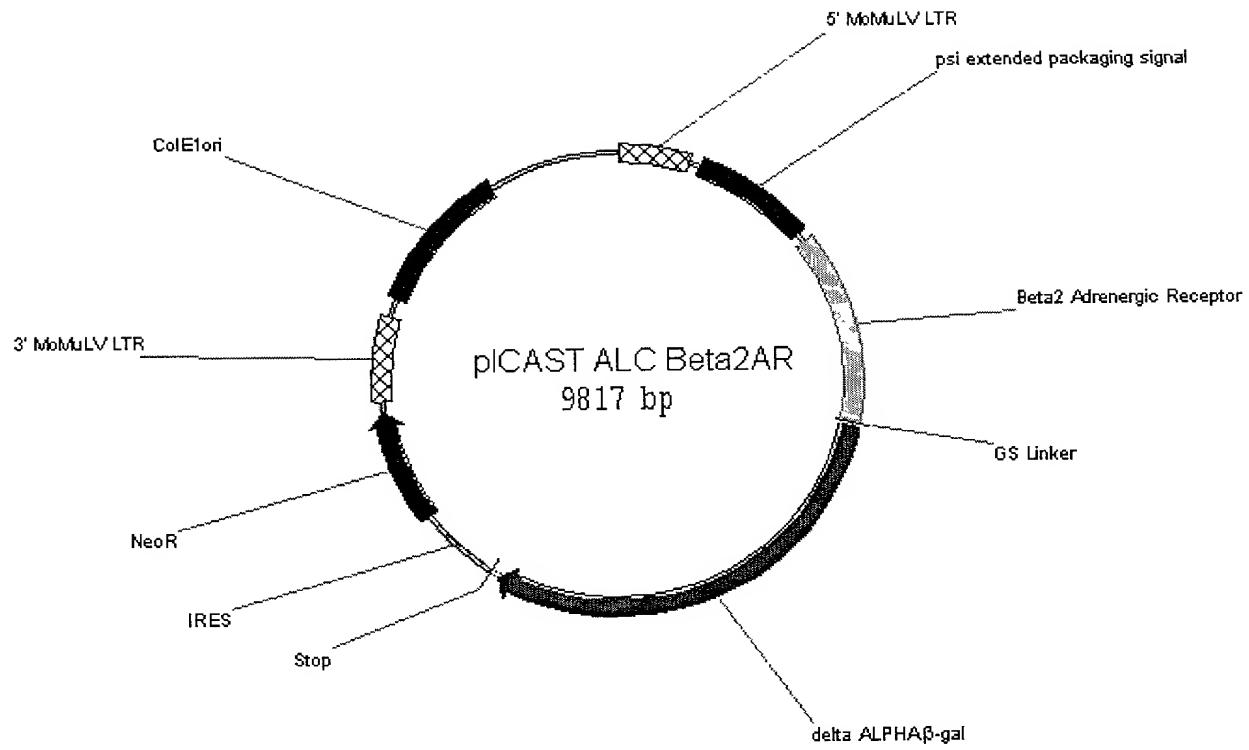


Figure 18

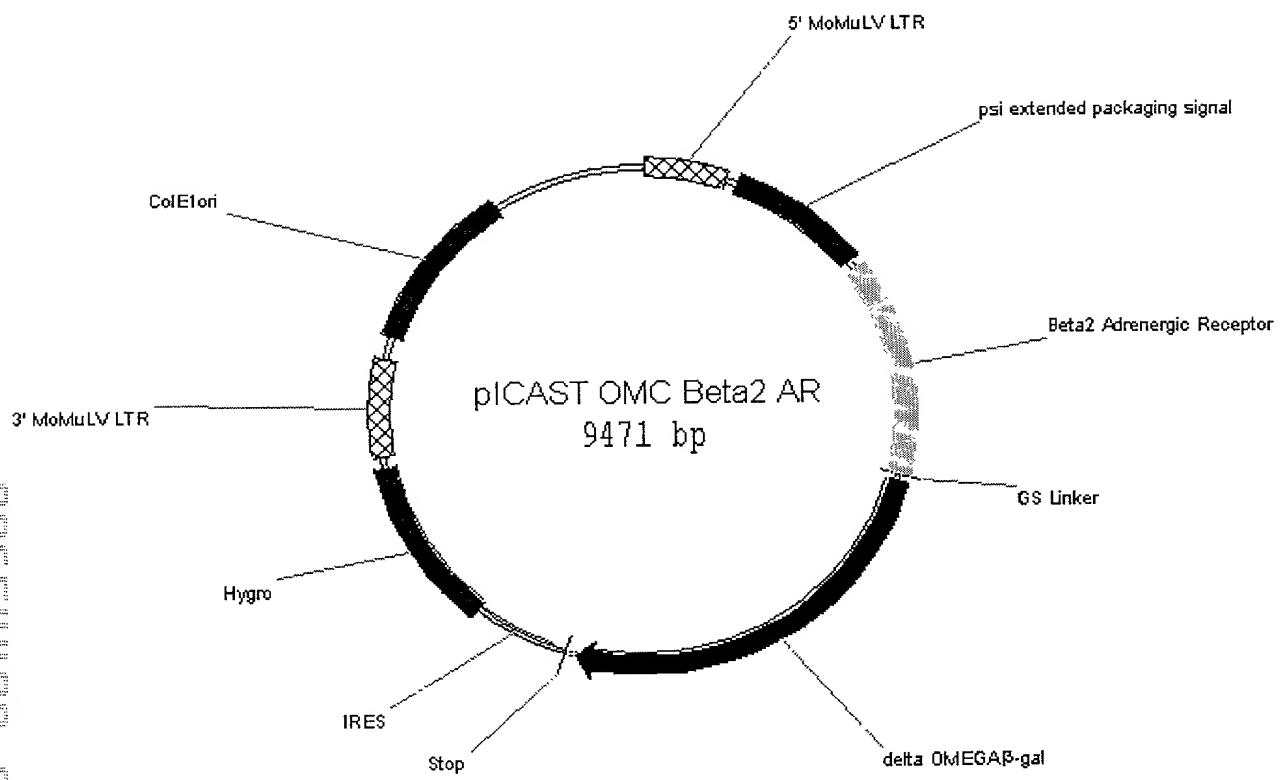


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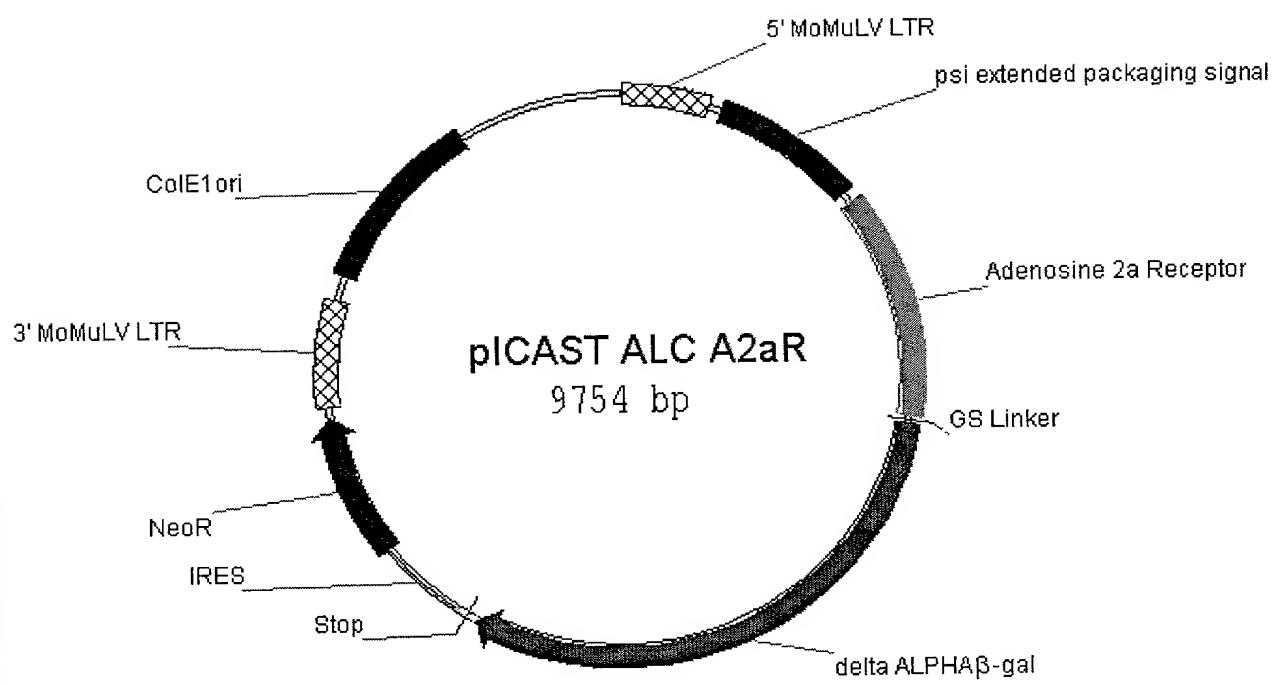


Figure 20

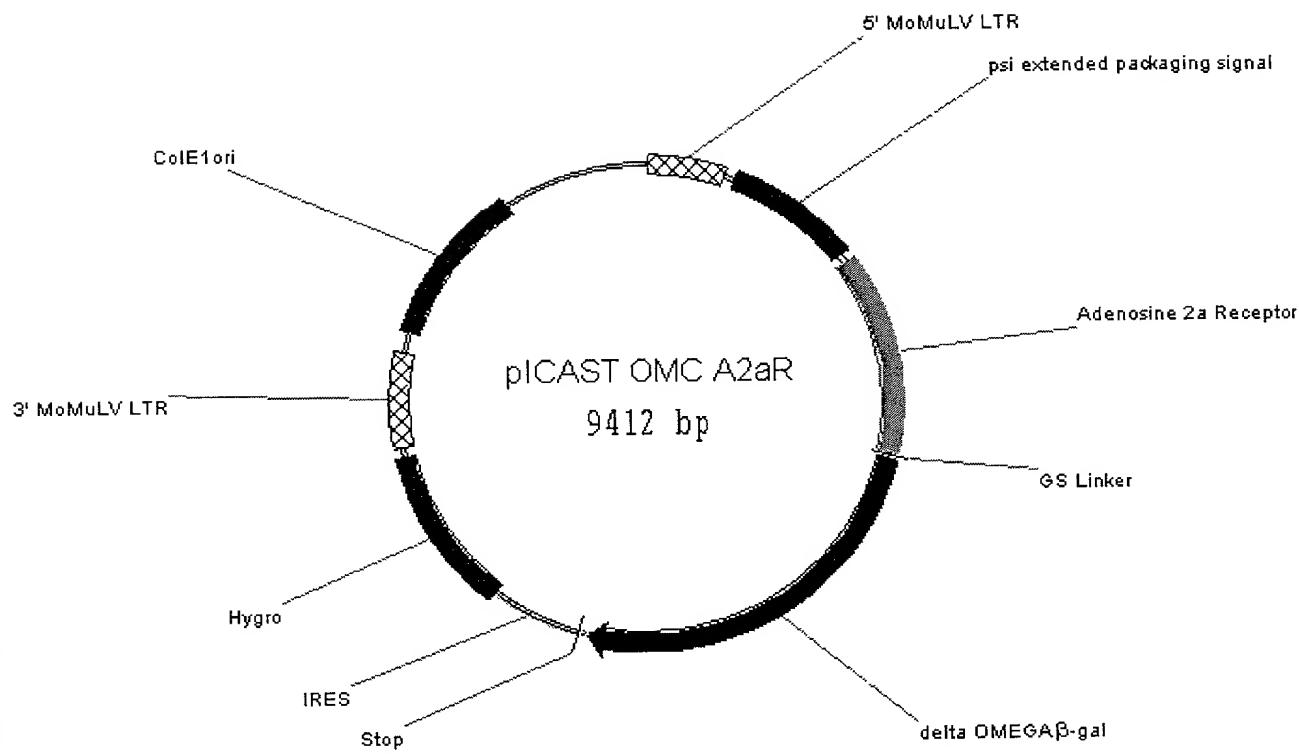


Figure 21

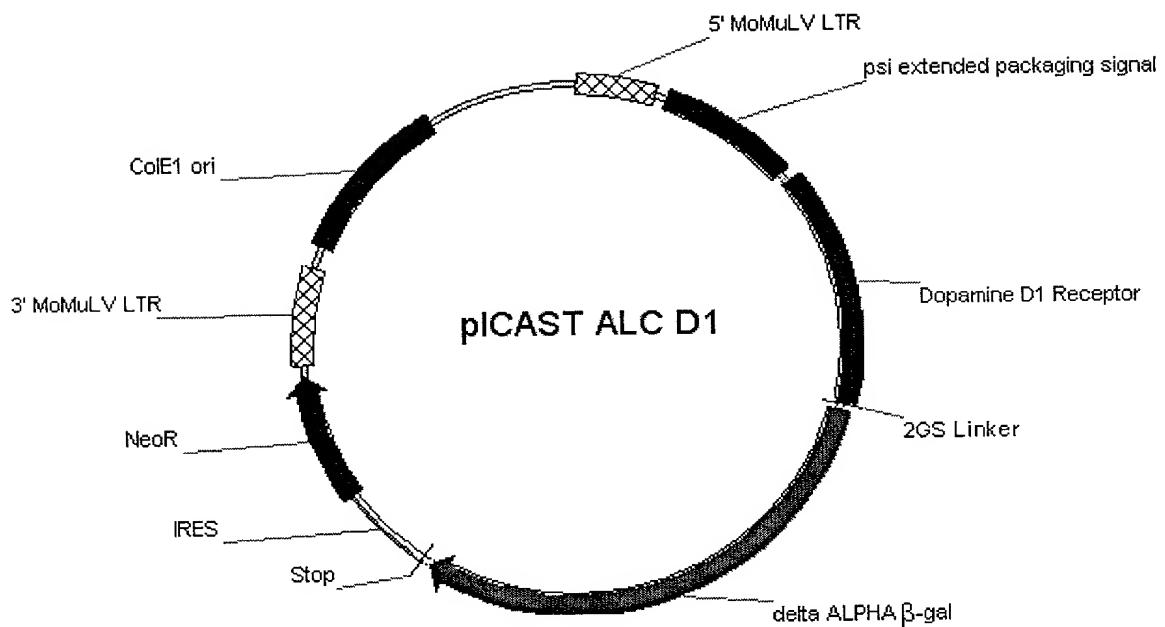
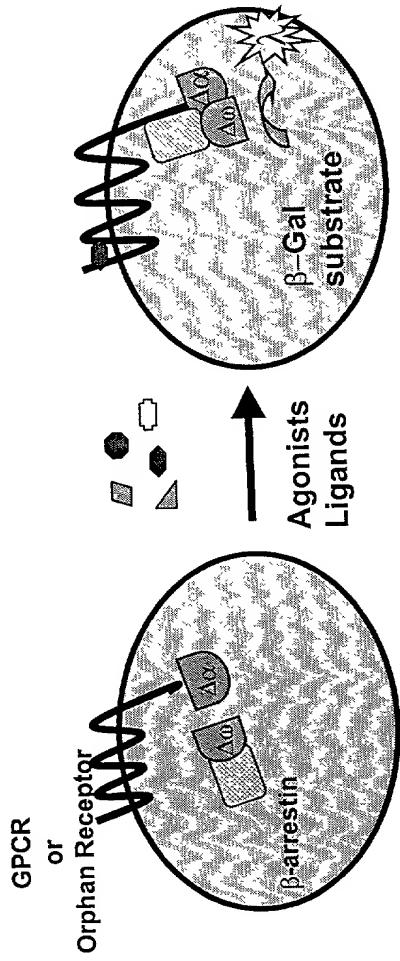


Figure 22

**Functional GPCR Activation Assay and Ligand Fishing for Orphan Receptors
by β -galactosidase mutant complementation in ICAST™ System**



Examples

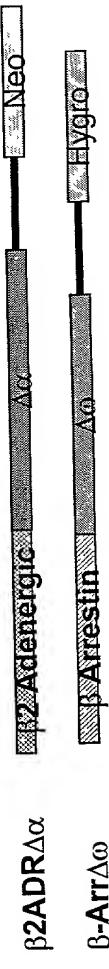


Figure 23

DOCKET NO. 4085-226-27

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF: Michelle A.J. PALMER, et al ART UNIT:
SERIAL NO.: New Application EXAMINER:
FILING DATE: Herewith
FOR: RECEPTOR FUNCTION ASSAY FOR G-PROTEIN COUPLED
RECEPTORS AND ORPHAN RECEPTORS BY REPORTER ENZYME
MUTANT COMPLEMENTATION

LIST OF INVENTORS' NAMES AND ADDRESSES

ASSISTANT COMMISSIONER FOR PATENTS
WASHINGTON, D.C. 20231

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A declaration containing all the necessary information will be submitted at a later date.

Respectfully submitted,

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